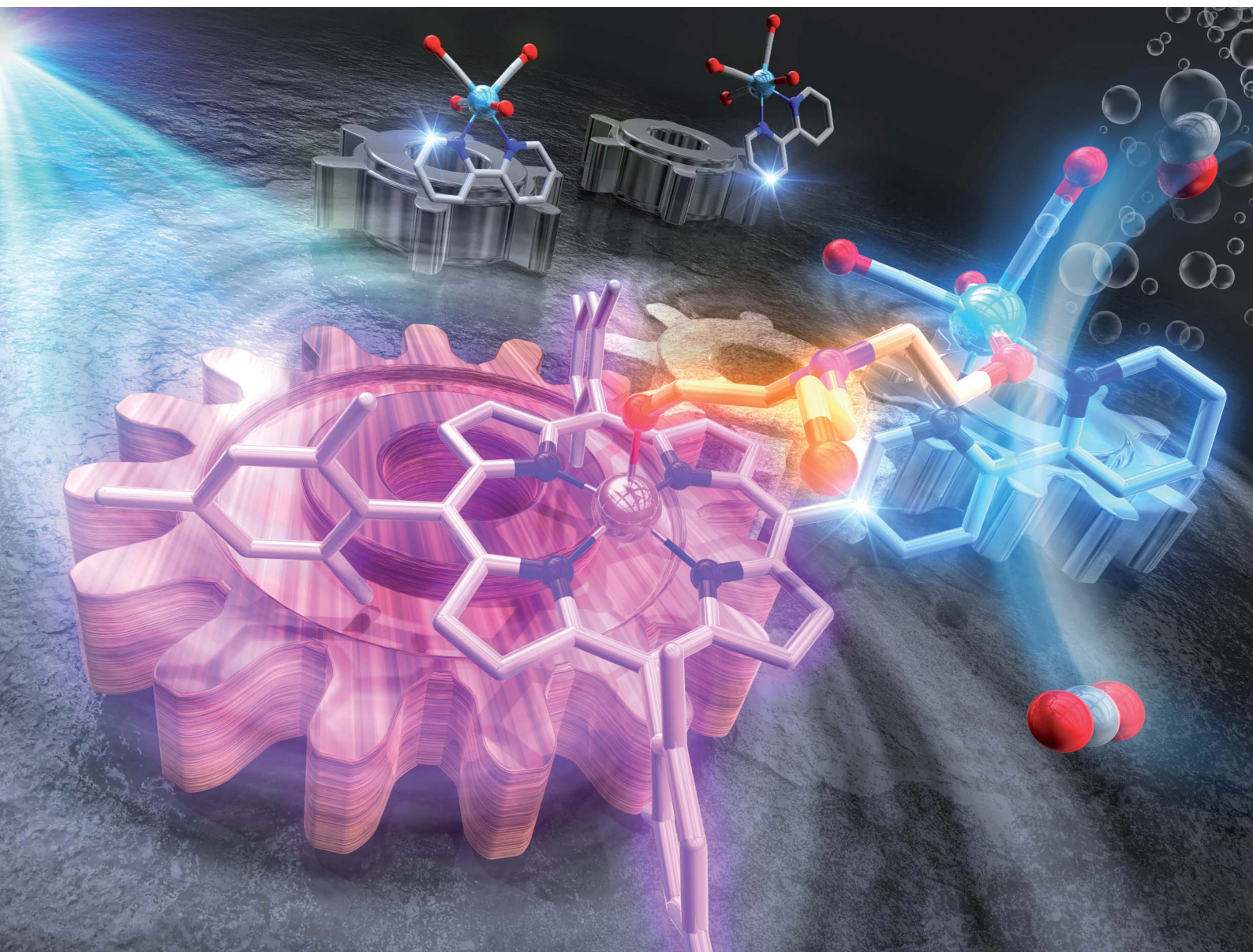


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EDGE ARTICLE

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Significance of the connecting position between Zn(II) porphyrin and Re(I) bipyridine tricarbonyl complex units in dyads for room-temperature phosphorescence and photocatalytic CO₂ reduction: unexpected enhancement by triethanolamine in catalytic activity†

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We synthesized three new dyads composed of a Zn porphyrin and *fac*-Re(bpy)(CO)₃Br (bpy = 2,2'-bipyridine) units, ZnP-*n*Bpy=Re^{Br} (*n* = 4, 5, and 6), in which the porphyrin is directly connected at the *meso*-position through the 4-, 5-, or 6-position of the bpy. We investigated the relationships between the connecting positions and the photophysical properties as well as catalytic activity in the CO₂ reduction reaction. The dyad connected through the 6-position, ZnP-6Bpy=Re^{Br}, showed obvious phosphorescence with a lifetime of 280 μs at room temperature, in *N,N*-dimethylacetamide (DMA), whereas the other two dyads showed almost no phosphorescence under the same conditions. The photocatalytic CO₂ reduction reactions in DMA using 1,3-dimethyl-2-phenyl-2,3-dihydro-1*H*-benzo[*d*]imidazole as the electron donor and the three dyads ZnP-*n*Bpy=Re^{Br} selectively produced CO with similar initial rates, but the durabilities were low. The addition of triethanolamine (TEOA) suppressed the decomposition of dyads, improving their durabilities and reaction efficiencies. In particular, ZnP-5Bpy=Re^{Br} was remarkably improved—it gave the highest durability and reaction efficiency among the three dyads; the reaction quantum yield reached 24%. The reason for this significant activity is no accumulation of electrons on the Zn porphyrin in ZnP-5Bpy=Re^{Br}, which would be caused by dual interactions of TEOA with the Re and Zn ions in the dyad. As the highest catalytic activity was observed in ZnP-5Bpy=Re^{Br} among the three dyads, which had no room-temperature phosphorescence (RTP), the catalytic activities and RTP properties are considered independent, but they are greatly influenced by the connecting positions on the bpy ligand in ZnP-*n*Bpy=Re^{Br}.

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Introduction

The global warming problem and fuel resource shortage necessitates research on artificial photosynthetic systems that use solar energy to reduce CO₂ to carbon resources. In constructing an artificial photosynthesis system, it is essential to

develop a photosensitizer that efficiently captures dilute sunlight,¹ and a catalyst that promotes the multielectron reduction reaction and suppresses H₂ evolution.² As a catalyst, metal complexes are promising candidates for CO₂ reduction due to their multiple accessible redox states, high activation energies against proton reduction, and various molecular design possibilities on the combination of appropriate metal ions and ligands. Among them, Re(I) diimine carbonyl complexes are advantageous as catalysts for CO₂ reduction because the Re complexes almost exclusively produce CO without the formation of H₂ even in aqueous solutions.^{2b} After the first report appeared describing the use of *fac*-Re(bpy)(CO)₃Cl (bpy = 2,2'-bipyridine) by Lehn and co-workers in the 1980s,³ solvents containing triethanolamine (TEOA) have often been used in many photocatalytic systems that employ Re complexes. While TEOA acts as a sacrificial reducing reagent, TEOA still acts as an effective additive when use is

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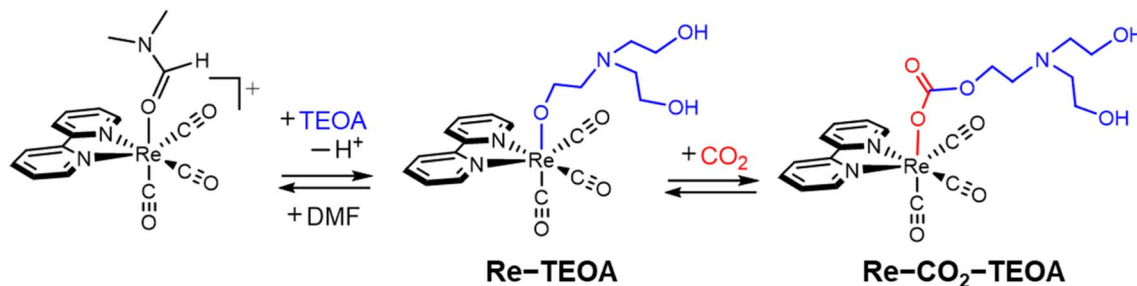


Fig. 1 Formation of the Re-CO₂-TEOA adduct through the Re-TEOA adduct in a CO₂-saturated DMF-TEOA solution.^{5a}

made of stronger and more efficient reducing reagents, where TEOA does then not act as a reducing reagent. In particular, the mixed solvent of DMF-TEOA (DMF = *N,N*-dimethylformamide) or DMA-TEOA (DMA = *N,N*-dimethylacetamide)⁴ behaves as a “magic solvent” for photocatalytic CO₂ reduction when employing Re complexes. Ishitani and co-workers found that $\text{fac}[\text{Re}(\text{bpy})(\text{CO})_3(\text{DMF})]^+$ is converted to the TEOA-CO₂ adduct (Re-CO₂-TEOA) through the TEOA adduct (Re-TEOA) in DMF-TEOA under a CO₂ atmosphere (Fig. 1).⁵ Thus, CO₂ is efficiently captured by the Re complex, with the assistance of TEOA, facilitating the photocatalytic CO₂ reduction even under low-concentration CO₂.^{5a} The similar insertion reactions of CO₂ into the M-OR (R = alkyl or aryl groups) bonds have been observed⁶ not only in Re complexes⁷ but also in the various other metal complexes,^{8–11} and are expected to be utilized in a wide range of fields such as CO₂ storage/release as well as CO₂ reduction reaction.

The Re complex also acts as a photosensitizer because it exhibits absorption in the near-ultraviolet to visible region; however, its durability is low. On the other hand, combination with a suitable photosensitizer, such as a Ru(II) diimine complex ($[\text{Ru}(\text{N}^{\wedge}\text{N})_3]^{2+}$), improves the efficiency and durability of photocatalytic CO₂ reduction.^{2b} The Ru complex has relatively strong absorption bands around 450 nm and can efficiently drive photoinduced electron transfer reactions by excitation with visible light. When a Ru diimine complex was connected with a Re complex (a Ru-Re complex), the photocatalytic CO₂ reduction selectively produced CO in the presence of 1,3-dimethyl-2-phenyl-2,3-dihydro-1*H*-benzo[*d*]-imidazole (BIH) as an electron donor—the turnover number (TON_{CO}) was > 2000 and the reaction quantum yield (Φ_{CO}) was 45%.¹² Furthermore, when the Ru-Re complex was combined with a semiconductor photocatalyst, then two-step photoexcitation of the Ru complex and the semiconductor selectively produced CO from CO₂ using water as the electron donor.¹³ The Ru diimine complexes have been utilized as the photosensitizer by combining other metal complex catalysts, showing high catalytic activity and durability in not only CO₂ reduction reaction but also oxygen and hydrogen evolutions from water.¹⁴ However, the absorption ability of the Ru diimine complex was insufficient for the full utilization of sunlight on the semiconductors.¹⁵ Therefore, the accumulation of photosensitizers¹⁶ and/or the development of photosensitizers that absorb visible light more strongly are essential.

In natural photosynthetic systems, networks of chlorophylls efficiently capture dilute photons from the sun and utilize the absorbed solar energy to produce chemical energy.¹⁷ Zn porphyrins are chlorophyll analogues—they have strong absorption bands in the visible region—and are used as photosensitizers in the photocatalytic CO₂ reduction.^{18–20} However, the use of Zn porphyrins as photosensitizers is often limited due to their poor stability under the photocatalytic reduction conditions because the accumulation of more than two electrons on the Zn porphyrins leads to hydrogenation, which finally results in color bleaching.^{19c,21}

Recently, we reported that photocatalytic CO₂ reduction using a dyad (ZnP-phen=Re, Fig. 2a) in which $\text{fac-Re}(\text{phen})(\text{CO})_3\text{Br}$ (phen = 1,10-phenanthroline) is directly linked to Zn tris(4-*tert*-butylphenyl)porphyrin selectively produced CO with a high reaction quantum yield ($\Phi_{\text{CO}} = 8\%$) and a high durability (TON_{CO} > 1400) in the presence of phenol as proton source and BIH as electron donor.²² The high durability is achieved by efficient electron transfer from the porphyrin to the adjacent Re complex where electrons are consumed by CO₂ reduction, thus suppressing the formation of two-electron reduced porphyrin that causes hydrogenation of the C=C bonds of the porphyrin skeleton. The high reaction quantum yield would be contributed to by the initial electron transfer from BIH through the long-lived excited triplet state (T₁) of porphyrin, enabling the excited species to be fully utilized in the electron transfer.^{22a} The efficient electron transfer from the T₁ is evidenced by the fact that the room-temperature phosphorescence (RTP) of the Zn porphyrin unit is remarkably quenched by a trace amount of BIH. In general, the phosphorescence from Zn porphyrin is not observed at room temperature, but only at low temperature. Thus, the unique RTP from the Zn porphyrin unit in ZnP-phen=Re would be imposed by the spin-orbital coupling by the heavy atoms of the Re complex.

In another paper, we reported that a slipped-cofacial porphyrin dimer (ReD', Fig. 2b) mimicking the natural special pair in photosynthesis exhibits high catalytic activity even under irradiation with strong light.²³ This is attributed to the intramolecular electron transfer from the lowest excited singlet state (S₁) of the porphyrin dimer to the Re complex unit, rather than the electron transfer through the T₁ of the porphyrin, resulting in no accumulation of the one-electron reduced species (OERS) on the porphyrin that causes the inner-filter effect.^{24,25} However, ReD' requires a high concentration of BIH, yet the reaction



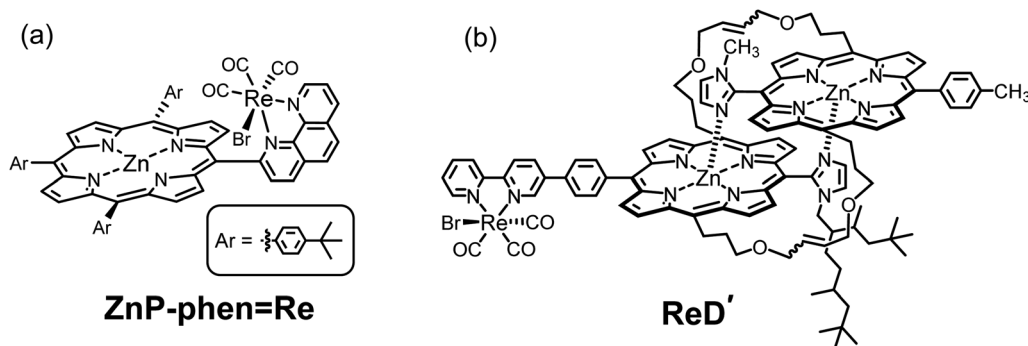


Fig. 2 Structures of (a) a previously reported dyad (ZnP-phen=Re)²² and (b) the special-pair mimic porphyrin dimer (ReD')²³

quantum yield is still lower (2%) than that of ZnP-phen=Re under weak light irradiation because a rapid back-electron transfer process generally competes with the catalytic reaction in the intramolecular charge-separated state.

ZnP-phen=Re exhibits unique properties: RTP from the Zn porphyrin and excellent activity for photocatalytic CO₂ reduction. However, it is unknown how the spatial distance and connecting positions between the Zn porphyrin and the Re complex units affect the RTP and the photocatalytic activity. Although the phen ligand was reported to show interesting properties, systematic modification of the phen part does not seem to be easy. The bpy ligand is used instead of the phen because the structure of bpy can be easily synthesized from a pyridyl precursor by a Pd-catalyzed coupling reaction, and various substituents can be introduced at the bpy part. Furthermore, in a *fac*-Re(diimine)(CO)₃Br-type complex the bpy ligand exhibits similar photocatalytic activity for CO₂ reduction to the phen ligand.²⁶

We have synthesized three new dyads, ZnP-*n*Bpy=Re^{Br} (*n* = 4, 5, and 6), in which *fac*-Re(bpy)(CO)₃Br (bpy = 2,2'-bipyridine) is directly connected with Zn(II) tris(2,4,6-trimethylphenyl) porphyrin at the *meso*-position through the 4, 5, or 6-position of the bpy (Fig. 3). The photochemical properties of ZnP-*n*Bpy=Re^{Br} were investigated, including RTP and photocatalytic activity for CO₂ reduction. When using ZnP-*n*Bpy=Re^{Br} in DMA

containing BIH as the electron donor, the photocatalytic CO₂ reduction had comparable initial production rates of CO as achieved with ZnP-phen=Re. The catalytic activities of ZnP-*n*Bpy=Re^{Br} were remarkably enhanced in the presence of TEOA, whereas ZnP-phen=Re was rather deactivated with TEOA. The reaction quantum yield reached the high value of Φ_{CO} = 24% in ZnP-5Bpy=Re^{Br}; it showed little RTP. The reaction quantum yield of ZnP-5Bpy=Re^{Br} was more than twice that of the other dyads. ZnP-5Bpy=Re^{Br} suppressed the electron accumulation at the porphyrin unit during the photocatalytic CO₂ reduction with unexpected specific assistance of TEOA.

Results and discussion

Synthesis

The synthetic route is shown in Scheme 1. Porphyrins in which a bpy group is directly connected at the *meso*-position have been synthesized by various procedures.^{27–29} We selected a synthetic route in which the bromopyridyl-substituted porphyrins are first synthesized and then a pyridyl group is introduced by reaction with the porphyrins using Pd-catalyzed Migita–Kosugi–Stille coupling.³⁰ Three types of bromopyridyl-substituted porphyrins were synthesized by the Lindsey method³¹ in the presence of pyrrole and 2,4,6-trimethylbenzaldehyde using 2-bromo-pyridine derivatives with a formyl group at the 4-, 5-, or 6-position. Zinc metalation of the free-base porphyrins was performed prior to the subsequent coupling reaction to prevent coordination of the palladium ion to the porphyrin center. The coupling reaction using 2-(tributylstannyl)pyridine in the presence of Pd(PPh₃)₄ followed by purification with silica gel and anhydrous K₂CO₃/flash silica gel (1:9 w/w)³² afforded pure porphyrins with a bipyridyl group, ZnP-*n*Bpy. The reaction of ZnP-*n*Bpy with one equivalent of Re(CO)₅Br followed by recrystallization from CHCl₃/ethanol afforded ZnP-*n*Bpy=Re^{Br}.

All signals of the ¹H NMR spectra were assigned using 2D NMR (¹H–¹H correlation spectroscopy (COSY)). Judging from the three CO signals in ¹³C NMR spectra (Fig. S41, S46, and S51†) and the patterns of the CO stretching bands in the infrared (IR) spectra (KBr pellets) (Fig. S44, S49, and S54†),³³ the facial Re complexes are obtained as shown in Fig. 3.

We also prepared ZnP-5Bpy=Re^{MeCN} and ZnP-6Bpy=Re^{MeCN} by replacing the bromide ion with acetonitrile using

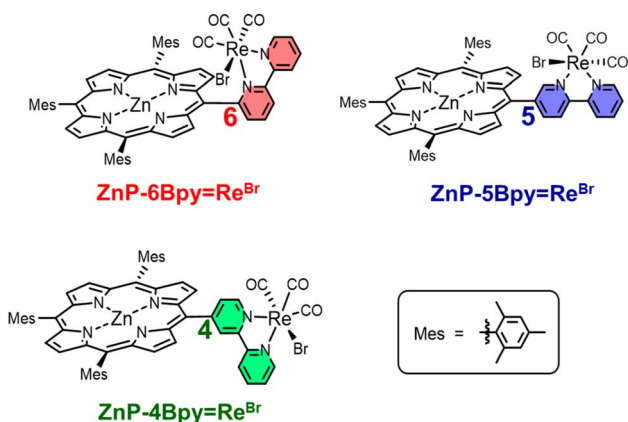
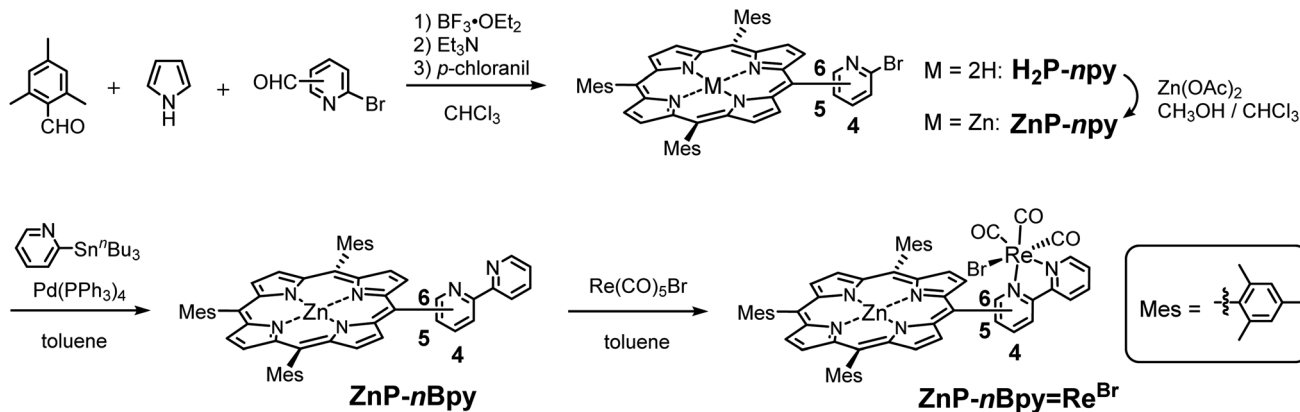
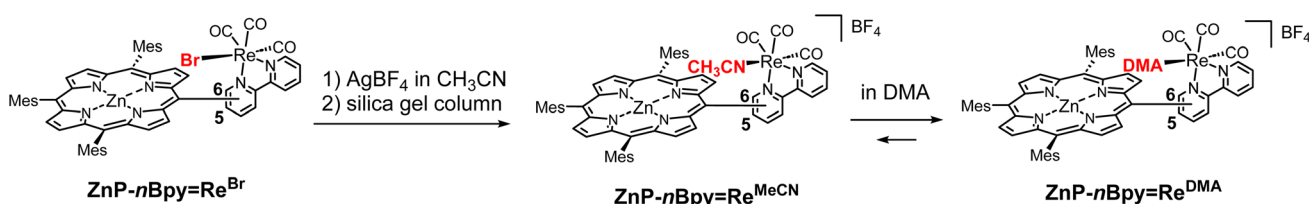


Fig. 3 Structures of three dyads, ZnP-*n*Bpy=Re^{Br} (*n* = 4, 5, and 6).



Scheme 1 Synthetic routes of $\text{ZnP-nBpy=Re}^{\text{Br}}$ ($n = 4, 5,$ and 6).Scheme 2 Synthesis of $\text{ZnP-nBpy=Re}^{\text{MeCN}}$ and ligand substitution of acetonitrile to DMA.

silver ions in the Re complex unit (Scheme 2). The substitutions of acetonitrile ligands with DMA in DMA solutions were confirmed by the observed changes in the stretching vibrations of CO in the IR spectra (Fig. S65[†]).^{5a}

Photophysical properties

The UV-vis absorption spectra of the dyads in DMA show intense Soret bands at approximately 430 nm and Q-bands at

>540 nm (Fig. 4a). Upon comparing the spectra of ZnP-nBpy and $\text{ZnP-nBpy=Re}^{\text{Br}}$, we note that the introduction of the Re tricarbonyl complex broadens the Soret bands and slightly red-shifts the Q-bands (Table 1 and Fig. S66[†]). This behavior was also observed in ZnP-phen=Re . Judging from the largest full width at half maxima (FWHM) of the Soret bands (Fig. 4a inset and Table 1), the electronic interaction of $\text{ZnP-6Bpy=Re}^{\text{Br}}$ is stronger (FWHM = 24 nm) than that of $\text{ZnP-4Bpy=Re}^{\text{Br}}$ (FWHM

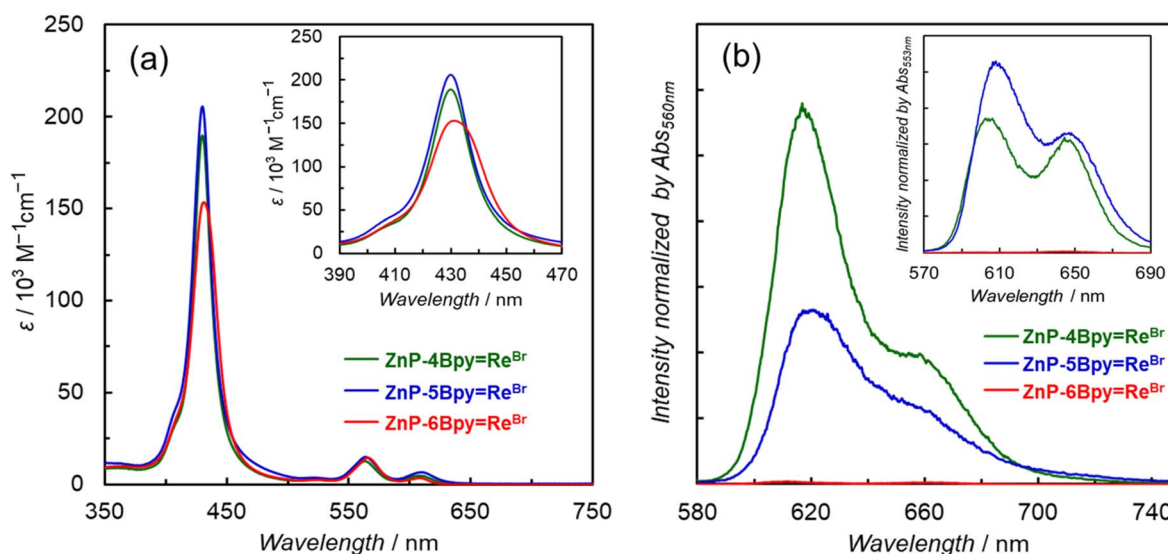


Fig. 4 (a) UV-vis absorption spectra of $\text{ZnP-nBpy=Re}^{\text{Br}}$ ($n = 4, 5,$ and 6) in DMA. Inset shows magnification of the Soret bands. (b) Fluorescence spectra of $\text{ZnP-nBpy=Re}^{\text{Br}}$ ($n = 4, 5,$ and 6) in Ar-saturated DMA ($\lambda_{\text{ex}} = 560$ nm) at 298 K. Inset shows spectra in Ar-saturated toluene ($\lambda_{\text{ex}} = 553$ nm) at 298 K. The fluorescence spectra were normalized with absorbance at the excitation wavelength.



Table 1 Absorption data in DMA, and fluorescence data in toluene and DMA^a

	$\lambda_{\text{abs}}/\text{nm}$ ($\epsilon/10^4 \text{ M}^{-1} \text{ cm}^{-1}$)	FWHM/nm	$\lambda_{\text{em}}^b/\text{nm}$	Φ_f^c	
				DMA	Toluene
ZnP-4Bpy	428 (50), 561 (1.7), 600 (0.52)	7.8	606	3.6%	3.3%
ZnP-4Bpy=Re ^{Br}	430 (19), 563 (1.3), 608 (0.45)	17	617	2.0% (0.56)	1.8% (0.55)
ZnP-5Bpy	429 (44), 562 (1.7), 602 (0.65)	9.4	605	2.6%	3.6%
ZnP-5Bpy=Re ^{Br}	430 (21), 564 (1.5), 609 (0.66)	18	621	1.1% (0.42)	2.2% (0.61)
ZnP-6Bpy	428 (42), 561 (1.5), 600 (0.44)	8.2	605	4.2%	3.6%
ZnP-6Bpy=Re ^{Br}	431 (15), 566 (1.5), 608 (0.36)	24	611	~0.02% (~0.004)	~0.03% (~0.009)
ZnP-phen=Re	430 (20), 565 (2.1), 606 (0.89)	29	610	~0.06%	~0.04% (~0.01)

^a The solutions were degassed by bubbling with Ar gas. Excited at 560 nm in DMA and 553 nm in toluene. ^b In DMA. ^c The values in parentheses are the ratio to the fluorescence quantum yield of the corresponding porphyrin before the introduction of the Re complex.

= 17 nm) and ZnP-5Bpy=Re^{Br} (FWHM = 18 nm). On the other hand, broadening of the absorption band occurs regardless of the distance between the Zn and Re atoms (Fig. S67[†]), suggesting that the electronic interaction is mediated *via* the bipyridine moieties. The solvent-substituted dyads, ZnP-5Bpy=Re^{DMA} and ZnP-6Bpy=Re^{DMA}, exhibited nearly identical absorption spectrum shapes to their corresponding Br-substituted dyads (Fig. S68[†]). The charge of the Re complex unit seems to have little effect on the absorption of the porphyrin unit.

Fluorescence spectra of the dyads are shown in Fig. 4b. Herein, DMA can coordinate to the central Zn ion of the porphyrin, affecting the spectral shape and wavelength.^{22b} The fluorescence of the porphyrin is almost completely quenched in ZnP-6Bpy=Re^{Br}, while the fluorescence quenching due to the introduction of the Re complex is moderate in ZnP-4Bpy=Re^{Br} and ZnP-5Bpy=Re^{Br} (Table 1 and Fig. S69[†]). The fluorescence lifetimes were measured in polar DMA (Fig. S70–S73[†]) and less polar toluene, to estimate the radiative and nonradiative rate constants together with the fluorescence quantum yields (Table 2). Upon comparing ZnP-6Bpy with ZnP-6Bpy=Re^{Br}, we note that the nonradiative rate constants are remarkably enhanced by the introduction of the Re complex regardless of the solvent ($k_{\text{nr}} = 3.8 \times 10^8 \text{ s}^{-1}$ for ZnP-6Bpy and $\sim 1 \times 10^{11} \text{ s}^{-1}$ for ZnP-6Bpy=Re^{Br} in DMA), suggesting that there is a rapid intersystem crossing (ISC) induced by the Re complex. A large k_{nr} value was also observed in ZnP-phen=Re. In ZnP-4Bpy=Re^{Br}

and ZnP-5Bpy=Re^{Br}, the nonradiative rate constants were also enhanced by introduction of the Re complex, although the extent of enhancement is not as great as in ZnP-phen=Re and ZnP-6Bpy=Re^{Br}. In all the dyads, the k_{nr} values are independent of the solvent polarity, indicating that the main nonradiative process from the S₁ of the porphyrin is the ISC on the porphyrin, which is imposed by the Re complex, rather than the photoinduced electron transfer from the S₁ of the Zn porphyrin unit to the Re complex unit as observed in ReD' (Fig. 2).²³ The nonradiative rate constant from the S₁ of the porphyrin of ZnP-6Bpy=Re^{Br} is approximately two orders of magnitude greater than for the other two dyads (1.3×10^9 and $1.5 \times 10^9 \text{ s}^{-1}$ for ZnP-4Bpy=Re^{Br} and ZnP-5Bpy=Re^{Br} in DMA, respectively), suggesting that the Zn porphyrin connected through the 6-position of the bpy is significantly influenced by spin-orbit coupling by the Re complex, whereas the porphyrins connected through the 4- and 5-positions are similarly and moderately affected.

ZnP-phen=Re showed phosphorescence from the Zn porphyrin in Ar-saturated DMA even at room temperature;^{22a} ZnP-6Bpy=Re^{Br} showed similar RTP. Fig. 5a shows the emission spectra at room temperature with delay time > 90 μs after the excitation pulse. The emission lifetimes were estimated to be 230 and 280 μs for ZnP-phen=Re and ZnP-6Bpy=Re^{Br}, respectively (Fig. 5b). Observation of the phosphorescence spectrum for Zn porphyrin at room temperature is extremely rare in solution.

Table 2 Fluorescence lifetimes (τ_f), radiative (k_f) and nonradiative rate constants (k_{nr}) from the S₁ of porphyrin at room temperature^{a,b}

	In DMA			In toluene		
	τ_f/ns	$k_f/10^9 \text{ s}^{-1}$	$k_{\text{nr}}/10^9 \text{ s}^{-1}$	τ_f/ns	$k_f/10^9 \text{ s}^{-1}$	$k_{\text{nr}}/10^9 \text{ s}^{-1}$
ZnP-4Bpy	2.2	0.016	0.44	2.1	0.016	0.46
ZnP-4Bpy=Re ^{Br}	0.76	0.026	1.3	0.57	0.032	1.7
ZnP-5Bpy	2.5	0.010	0.39	2.5	0.014	0.39
ZnP-5Bpy=Re ^{Br}	0.68	0.016	1.5	0.67	0.033	1.5
ZnP-6Bpy	2.5	0.017	0.38	2.5	0.014	0.39
ZnP-6Bpy=Re ^{Br}	~0.01	~0.02	~100	~0.01	~0.03	~100
ZnP-phen=Re	~0.02	~0.03	~50	~0.02	~0.03	~70

^a Excited at 406 nm (FWHM 70 ps) in DMA and in toluene. ^b Radiative rate constant: $k_f = \Phi_f/\tau_f$. Nonradiative rate constant: $k_{\text{nr}} = 1/\tau_f - k_f$.



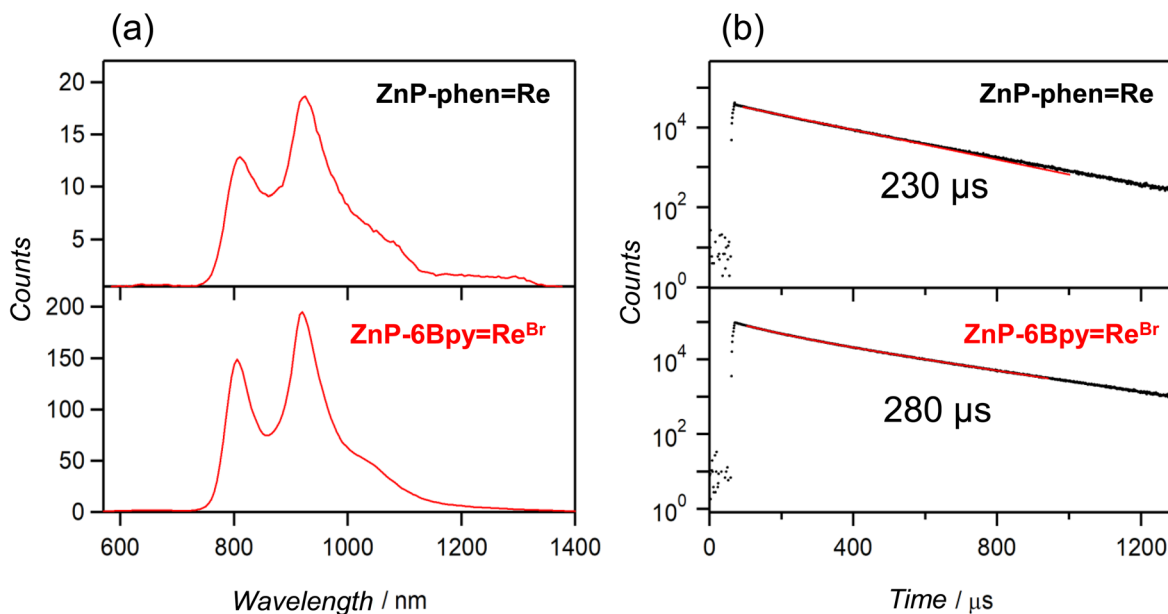


Fig. 5 (a) Phosphorescence spectra of ZnP-phen=Re and ZnP-6Bpy=Re^{Br} in Ar-saturated DMA ($\lambda_{\text{ex}} = 405 \text{ nm}$) at room temperature. The spectra were recorded with delay time $> 90 \mu\text{s}$ after the excitation pulse. (b) Phosphorescence decay profiles (semilog plot) of ZnP-phen=Re and ZnP-6Bpy=Re^{Br} in Ar-saturated DMA ($\lambda_{\text{ex}} = 405 \text{ nm}$) at room temperature, monitored at 800 and 920 nm. The decay profiles were fitted with one component (red lines).

In contrast to ZnP-6Bpy=Re^{Br}, ZnP-4Bpy=Re^{Br} showed very weak phosphorescence, while phosphorescence from ZnP-5Bpy=Re^{Br} was hardly detected in DMA (Fig. 6a and S74[†]). The phosphorescence lifetimes of ZnP-4Bpy=Re^{Br} and ZnP-5Bpy=Re^{Br} were preliminarily estimated to be several tens of microseconds—this is an order of magnitude smaller than for ZnP-6Bpy=Re^{Br}. Considering that almost no phosphorescence was observed, the shorter lifetimes would originate from competitive rapid thermal deactivation processes.

To investigate the electron transfer process through the T_1 of the porphyrin, quenching experiments with the addition of BIH were carried out. As observed in ZnP-phen=Re,^{22a} the phosphorescence of ZnP-6Bpy=Re^{Br} was efficiently quenched by the addition of BIH (Fig. 6b). The Stern–Volmer constant was estimated to be $K_{\text{SV}} = 11\,000 \text{ M}^{-1}$. On the other hand, compared with the value of ZnP-phen=Re ($K_{\text{SV}} = 180\,000 \text{ M}^{-1}$),^{22a} the value of $11\,000 \text{ M}^{-1}$ for ZnP-6Bpy=Re^{Br} is almost an order of magnitude smaller. Considering that the phosphorescence lifetimes of ZnP-phen=Re (230 μs) and ZnP-6Bpy=Re^{Br} (280 μs) are similar, the smaller Stern–Volmer constant in ZnP-6Bpy=Re^{Br} would come from the smaller quenching rate constants (k_q) due to the slower electron transfer rate from BIH to the excited porphyrin (*vide infra*).

We also performed emission measurements on ZnP-5Bpy=Re^{DMA} and ZnP-6Bpy=Re^{DMA} (Fig. S75[†]), showing that the fluorescence intensities remained unchanged, and in the case of ZnP-6Bpy=Re^{DMA}, the RTP was observed. This suggests that even in the dyads with the solvent-substituted cationic Re complexes, the intramolecular photoinduced electron transfer is a minor process, contrasting with previous studies^{19c,e,34,35} and observation in ReD'.²³ In our system, the porphyrin and Re

complex are in close proximity. This proximity effect may destabilize charge-separated states and lead to intersystem crossing. To clarify these details, further time-resolved absorption and infrared spectroscopic measurements are currently underway.

Electrochemical properties

Cyclic voltammograms (CVs) of the ZnP-*n*Bpy series show that first reduction waves are reversible in Ar-saturated DMA (Fig. S76,† dotted lines), indicating that the OERS of the Zn porphyrin is stable in DMA. The redox potentials obtained from the differential pulse voltammograms (DPVs) of the dyads and the corresponding porphyrins are summarized in Table S1.† The first reduction peaks of the ZnP-*n*Bpy=Re^{Br} series appear at -1.68 V ($n = 4$), -1.64 V ($n = 5$), and -1.76 V ($n = 6$), which are on the more positive side than the reduction peaks (-1.84 to -1.87 V) of the ZnP-*n*Bpy series. The first reduction potentials of the ZnP-*n*Bpy=Re^{Br} series were rather similar to that of *fac*-Re(bpy)(CO)₃Br (-1.73 V). Density functional theory (DFT) calculations with consideration of the solvent (Fig. S77[†]) indicate that the lowest unoccupied molecular orbitals (LUMOs) and spin density plots for the OERS are abundantly distributed on the Re complex (Fig. S78 and S79[†]), indicating that the first reductions of the dyads occur mainly on the Re complex units of the ZnP-*n*Bpy=Re^{Br} series in the electrochemical measurements. The reduction potentials of the ZnP-*n*Bpy series are more negative (-1.84 to -1.87 V) than that of the porphyrin unit of ZnP-phen=Re (-1.79 V).^{22a} The reduction potential of Zn *meso*-tetramesitylporphyrin (ZnTMP) is reported to be 180 mV more negative than that of Zn *meso*-tetraphenylporphyrin (ZnTPP), and the substituent effect of the *tert*-butyl groups is



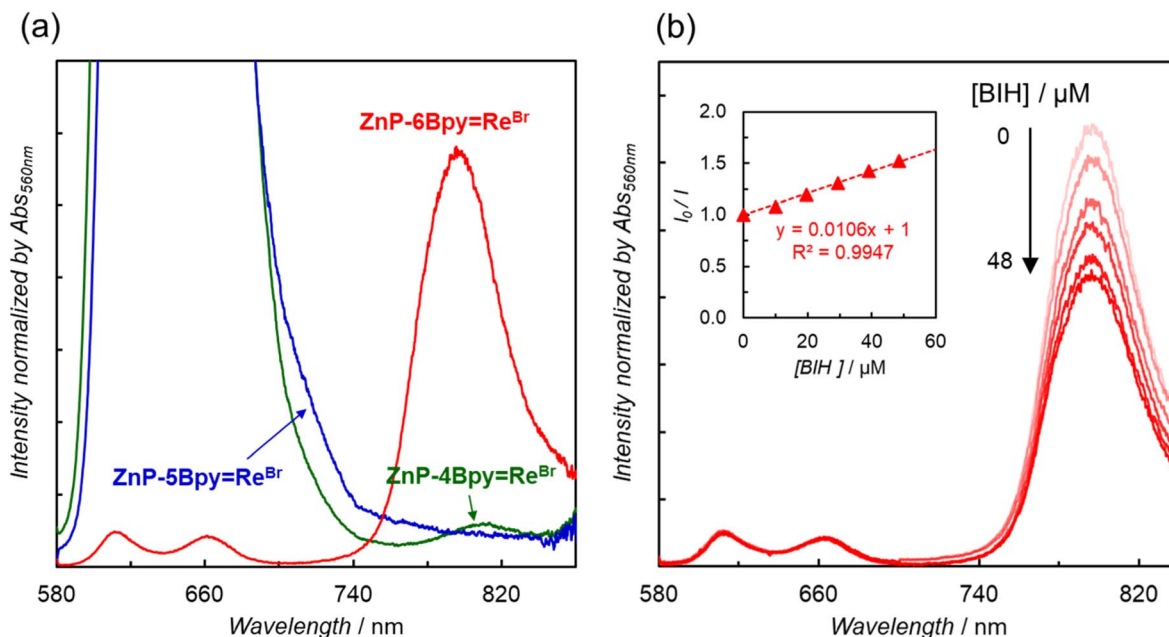


Fig. 6 (a) Comparison of emission spectra of ZnP-*n*Bpy=Re^{Br} in Ar-saturated DMA ($\lambda_{\text{ex}} = 560$ nm) at 298 K. (b) Emission spectra of ZnP-6Bpy=Re^{Br} in the presence of various amounts of BIH in Ar-saturated DMA ($\lambda_{\text{ex}} = 560$ nm) at 298 K. Inset shows Stern–Volmer plot of the emission.

small ($\Delta E^{\text{red}} = 20$ mV in Fig. S80c†).³⁶ Thus, the negative shifts arise from the substituents at the *meso*-positions of the porphyrin unit: the ZnP-*n*Bpy=Re^{Br} series has mesityl groups while the porphyrin unit of ZnP-phen=Re has 4-*tert*-butyl-phenyl groups.

Assuming that the energy levels of the T₁ of the ZnP-*n*Bpy=Re^{Br} series are similar to in ZnP-phen=Re and that the reduction potential of the porphyrin moiety remains the same before and after introduction of the Re complex moiety, the electron transfer from BIH to the T₁ is thermodynamically unfavorable (Fig. S81†). However, the phosphorescence quenching experiment gave a Stern–Volmer constant in ZnP-6Bpy=Re^{Br} that reached $K_{\text{SV}} = 11\,000\text{ M}^{-1}$; this value is much greater than values recorded for the well-used [Ru(N[^]N)₃]²⁺.¹² This originates from the long T₁ lifetime (τ_p) of the Zn porphyrin, which allows the unfavorable electron transfer process. The k_q value for ZnP-6Bpy=Re^{Br} ($3.9 \times 10^7\text{ M}^{-1}\text{ s}^{-1}$) was smaller than that for ZnP-phen=Re ($7.8 \times 10^8\text{ M}^{-1}\text{ s}^{-1}$). This would be attributed to the more negative shift of the reduction potential of the Zn porphyrin unit due to the mesityl substituents.³⁷

Photocatalytic CO₂ reduction in DMA and additives suppressing dissociation of the Re complex

Photocatalytic CO₂ reductions were carried out in DMA containing the dyad and BIH as the electron donor (Fig. 7). Here, 0.05 M of BIH was used to almost completely quench the T₁ (99.8% in ZnP-6Bpy=Re^{Br}). Irradiation of a CO₂-saturated DMA solution containing ZnP-phen=Re and BIH showed a linear increase in CO formation with high durability, as observed previously.^{22a} Almost no activity difference was observed between ZnP-4Bpy=Re^{Br}, ZnP-5Bpy=Re^{Br}, and ZnP-6Bpy=Re^{Br}. Although the three dyads showed similar initial reaction

rates for CO formation to that of ZnP-phen=Re, their activities were lost within 1 h and TON_{CO} was <20 even after 8 h.

In the emission quenching experiments, we found that the fluorescence of ZnP-6Bpy=Re^{Br} increased during irradiation in the presence of BIH under a CO₂ atmosphere (Fig. 8a). No increase in the fluorescence was observed in ZnP-phen=Re or in the absence of BIH. The rate of fluorescence increase in ZnP-6Bpy=Re^{Br} was slower under a CO₂ atmosphere than under an

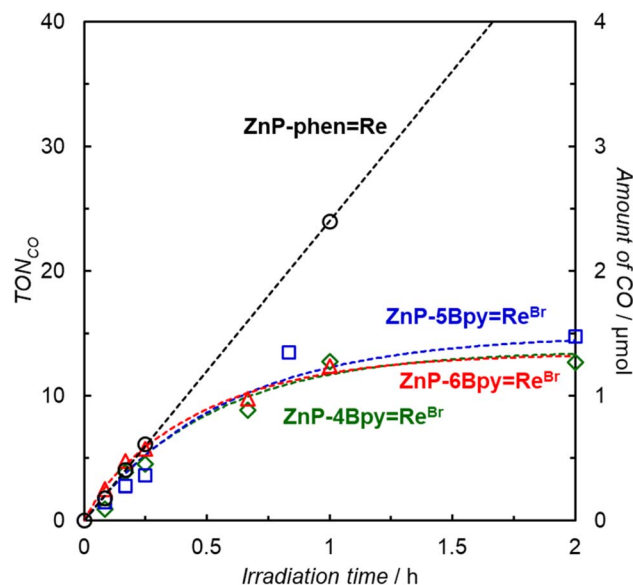


Fig. 7 Time dependence of CO formation (0–2 h) during irradiation at 420 nm by LED lamps (input power: 30 mW), in a merry-go-round irradiation apparatus, of CO₂-saturated DMA solutions containing the dyad (0.05 mM) and BIH (0.05 M).



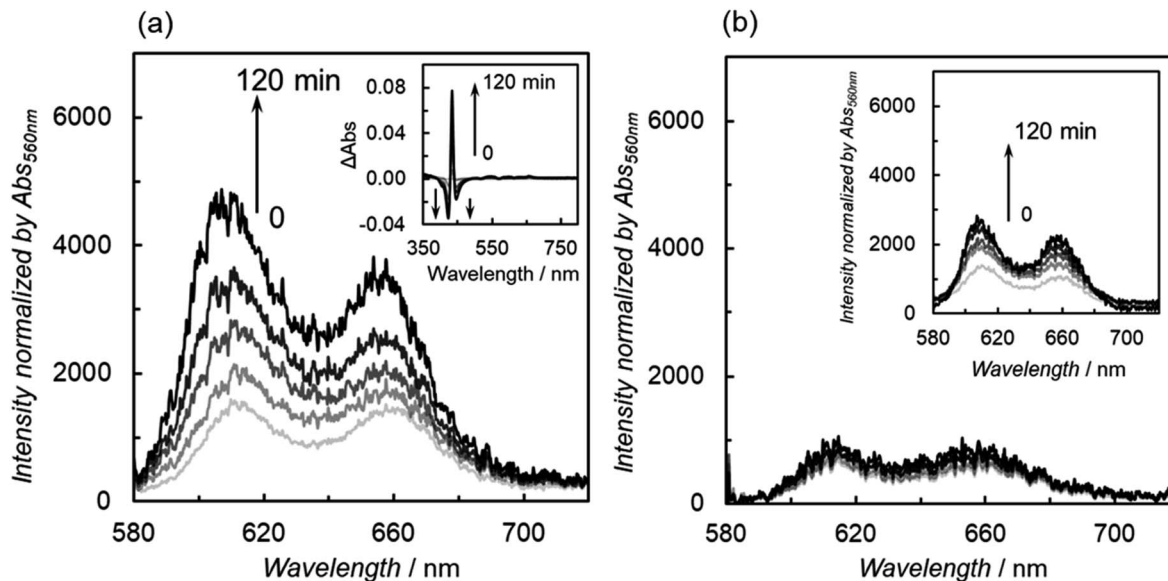


Fig. 8 (a) Fluorescence spectral changes (0–120 min) of the CO₂-saturated DMA containing ZnP-6Bpy=Re^{Br} (3 μM) in the presence of 50 μM of BIH during irradiation at 560 nm. Inset shows differential UV-vis absorption spectra during the irradiation. (b) Fluorescence spectral changes (0–120 min) of the CO₂-saturated DMA-TEOA (4 : 1 v/v) containing ZnP-6Bpy=Re^{Br} (3 μM) in the presence of 50 μM of BIH during irradiation at 560 nm. Inset shows fluorescence spectral change in the presence of 6.7 mM phenol.

Ar atmosphere. Differential UV-vis absorption spectra of ZnP-6Bpy=Re^{Br} during the irradiation show the appearance of a sharp peak in the Soret band corresponding to ZnP-6Bpy (Fig. 8a inset). Thus, it is thought that the reduced ZnP-6Bpy=Re^{Br} leads to dissociation of the Re tricarbonyl unit to revive the fluorescence quenched by the Re complex. The phen ligand of ZnP-phen=Re is structurally more rigid, with two nitrogen atoms fixed in the same direction, giving a more stable Re complex structure and high durability for the photocatalytic CO₂ reduction. Fig. 8b shows the fluorescence spectral changes in the presence of phenol and TEOA. Surprisingly, phenol partially suppresses dissociation of the Re tricarbonyl unit and TEOA almost completely suppresses the dissociation. The significant suppression of the decomposition by the additives was not observed under an Ar atmosphere. It has been reported that a proton promotes the reaction of the reduced Re complex with CO₂,³⁸ and TEOA coordinates to the Re complex in the ground state to form an adduct with CO₂.⁵ Otherwise, TEOA can reversibly react with CO₂ to form a zwitterionic adduct which acts as a good proton source (Fig. S82†).^{8a} Thus, TEOA might also contribute to the protonation of Re-COO⁻ to Re-COOH. In any case, the additives might change the electron distribution of the reduced Re complex upon reaction with CO₂, thus suppressing the dissociation of the Re tricarbonyl unit. Several deactivation pathways of Re diimine carbonyl complexes have been reported to involve the formation of Re formate complex,^{3a} Re dimeric species,³⁹ and so on.^{25a} To the best of our knowledge, the dissociation of the Re tricarbonyl moiety from the bpy ligand after reduction has not been reported as the catalytic deactivation pathway for the CO₂ reduction reactions. During the measurements of the emission spectra of the DMA-substituted cationic complex (ZnP-5Bpy=Re^{DMA} and ZnP-

6Bpy=Re^{DMA}), increases in fluorescence were observed under dilution conditions (Fig. S83†). On the other hand, no such spectral changes were observed with the Br-substituted complex (ZnP-5Bpy=Re^{Br} and ZnP-6Bpy=Re^{Br}). These results suggest that dissociation of Re ion occurs in the solvent-substituted cationic complex, lacking the Br ion. Since the fluorescence spectral changes were suppressed in the presence of TEOA (Fig. S84†), TEOA stabilizes the solvent-substituted species, probably by coordination to the cationic Re ion.

Photocatalytic CO₂ reduction in DMA-TEOA

Because the additives improved the stability of ZnP-6Bpy=Re^{Br}, the photocatalytic CO₂ reductions were carried out in the presence of phenol (Fig. S85†) and TEOA (Fig. 9). In particular, the addition of TEOA remarkably improved the catalytic activities of the ZnP-*n*Bpy=Re^{Br} series in terms of reaction rate and durability. Herein, the photocatalytic CO₂ reductions selectively produced CO without forming detectable amounts of CH₄ and HCOOH, but with a trace amount of H₂ (Fig. S86†). This improvement would result from the suppression of the Re dissociation by TEOA. Fig. 9 shows that only ZnP-phen=Re is deactivated in the presence of TEOA. Previously we reported that TEOA promotes the formation of chlorins and decreases the catalytic durability in ZnP-phen=Re.²³ Furthermore, the chlorins are formed by the reaction of TEOA with the excited porphyrin at the β-pyrrole position.⁴⁰ Thus, the lack of inactivation of the ZnP-*n*Bpy=Re^{Br} series would be attributed to the low reactivity of the porphyrins to TEOA, which would be caused by the mesityl groups having high steric hindrance and electron donating. Upon comparing the initial reaction rates, we note that all dyads in the ZnP-*n*Bpy=Re^{Br} series surpass the previous ZnP-phen=Re system under optimal conditions. In particular,



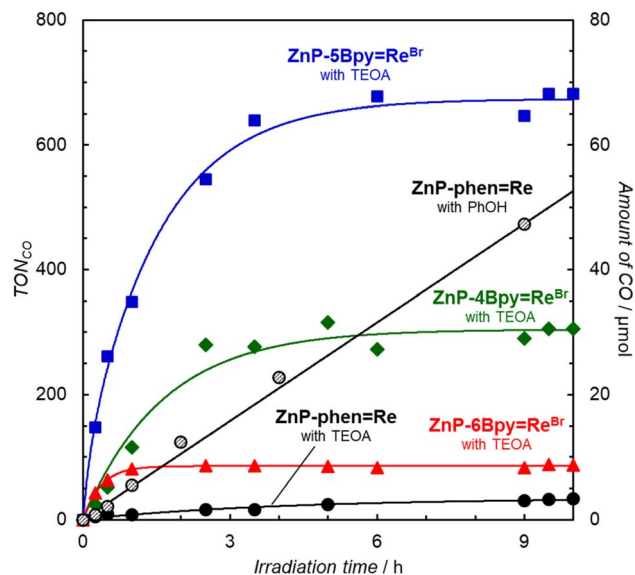


Fig. 9 Time dependence of CO formation during irradiation at 420 nm by LED lamps (input power: 30 mW), in a merry-go-round irradiation apparatus, of CO₂-saturated DMA-TEOA (4 : 1 v/v) solutions containing the dyad (0.05 mM) and BIH (0.05 M). ZnP-4Bpy=Re^{Br} (green-filled diamond), ZnP-5Bpy=Re^{Br} (blue-filled square), ZnP-6Bpy=Re^{Br} (red-filled triangle), ZnP-phen=Re (black-filled circle). Plot of ZnP-phen=Re (0.05 mM) in CO₂-saturated DMA containing BIH (0.05 M) and phenol (0.1 M) shown as black open circles.

ZnP-5Bpy=Re^{Br} unexpectedly exhibits a higher initial reaction rate (twice as high) and higher durability than achieved with ZnP-4Bpy=Re^{Br} and ZnP-6Bpy=Re^{Br}; the turn over frequency (TOF_{CO}) reaches 590 h⁻¹. However, even in ZnP-5Bpy=Re^{Br} the CO formation stopped within 5 h and further addition of BIH to the reaction mixture did not restart the photocatalytic reaction.

The UV-vis absorption spectra after 17 h showed degradation of the porphyrins (Fig. S87[†]), indicating that the photocatalysts decomposed during the irradiation. The UV-vis absorption spectral change of ZnP-5Bpy=Re^{Br} during relatively strong light irradiation at 420 nm showed the formations of the OERS of the porphyrin units at *ca.* 450 and 700 nm⁴¹ and chlorins at *ca.* 610 nm⁴² at the initial stage (Fig. S88[†]). Thus, the photocatalyst would undergo further reduction from chlorin and faded during the irradiation.

As it is thought that the accumulation of two electrons on the porphyrin causes the decomposition,²¹ we carried out the photocatalytic CO₂ reduction using weaker light to prevent the electron accumulation rate from exceeding the CO₂ reduction reaction rate. Fig. 10a shows the time-course comparison of CO production using ZnP-5Bpy=Re^{Br} when the input power of the apparatus was changed from 30 mW to 0.5 mW. As a control experiment, the reaction using a mixed system of ZnTMP and *fac*-Re(bpy)(CO)₃Br was carried out with weak light (0.5 mW). In the mixed system, CO production was mainly observed but the reaction stopped when the TON_{CO} reached approximately 100. On the other hand, in ZnP-5Bpy=Re^{Br} the CO production linearly increased to TON_{CO} ≈ 1000, corresponding to a quantitative amount of added BIH (1000 equivalents). Further addition of 500 equivalents of BIH restarted the reaction, with the same reaction rate, and the reaction continued until all BIH was consumed. A plot of TON_{CO} versus the product of input power and irradiation time shows that the reaction proceeds more efficiently with weak light irradiation (Fig. 10b).

Initial photoinduced electron transfer process from BIH

The photocatalytic CO₂ reduction using ZnP-5Bpy=Re^{Br} showed the highest reaction quantum yield with high

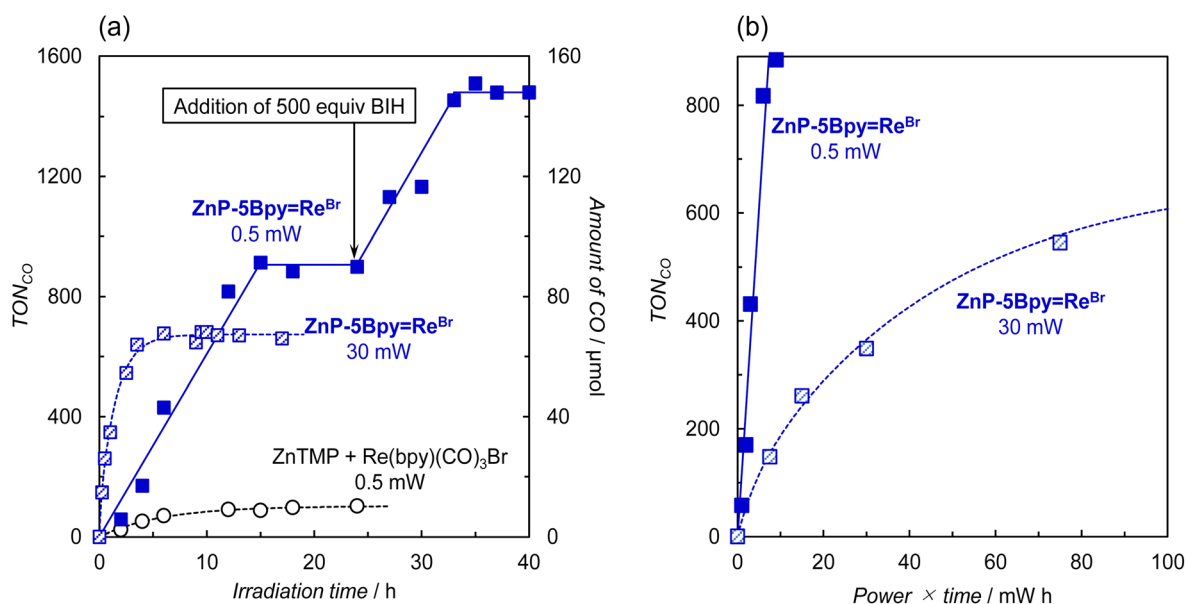


Fig. 10 (a) Time dependence of CO formation during irradiation at 420 nm by LED lamps (input power: 0.5 and 30 mW), in a merry-go-round irradiation apparatus, of CO₂-saturated DMA-TEOA (4 : 1 v/v) solutions containing ZnP-5Bpy=Re^{Br} (0.05 mM), and a mixed system of ZnTMP (0.05 mM) and *fac*-Re(bpy)(CO)₃Br (0.05 mM) in the presence of BIH (0.05 M). (b) Plots of TON_{CO} versus product of input power and irradiation time.



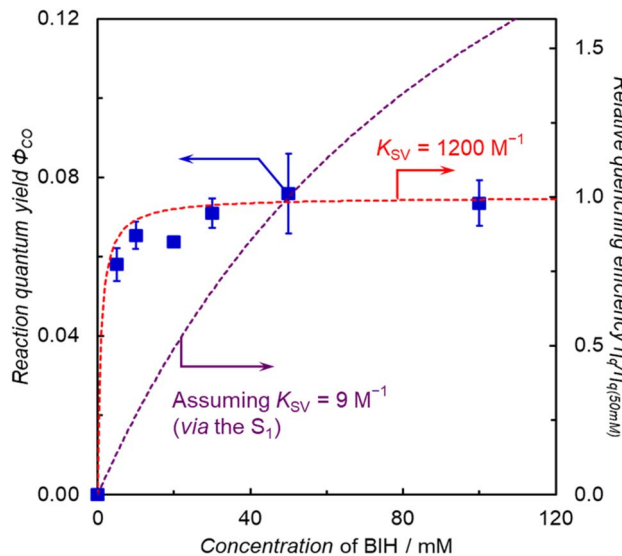


Fig. 11 Plot of the Φ_{CO} versus the initial concentration of BIH in the photocatalytic CO_2 reduction (Xe lamp, 420 nm, 3.6×10^{-8} einstein per s) using DMA-TEOA (4 : 1 v/v) solutions containing ZnP-5Bpy=Re^{Br} (0.05 mM). The quenching efficiency relative to η_q at [BIH] = 50 mM was calculated using $\eta_q = [\text{BIH}]/K_{\text{SV}}/(1 + [\text{BIH}]/K_{\text{SV}})$.

durability, but ZnP-5Bpy=Re^{Br} showed no RTP. This indicates that there is no direct correlation between the phosphorescence and the catalytic activity. On the other hand, the time course of CO production showed that the photocatalytic CO_2 reduction proceeded at a constant rate until all BIH in the system was consumed (Fig. 10a). Thus, the reaction proceeds efficiently even under dilute BIH conditions, indicating that the electron transfer from BIH proceeds through a long-lived excited state, that is, the T_1 of the porphyrin. In fact, the plots of Φ_{CO} of ZnP-5Bpy=Re^{Br} against the initial concentration of BIH (Fig. 11) show that the Φ_{CO} is almost constant and well fitted with the simulation curve assuming $K_{\text{SV}} = 1200 \text{ M}^{-1}$, where the value of k_q is similar to that of ZnP-6Bpy=Re^{Br} ($3.9 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$) and the phosphorescence lifetime assuming 30 μs . A similar independent plot of Φ_{CO} versus [BIH] is obtained for ZnP-phen=Re (Fig. S89[†]). Thus, in ZnP-5Bpy=Re^{Br}, even though no RTP was observed, the quantitative electron transfer from BIH to the T_1 of the porphyrin occurs, allowing the efficient photocatalytic CO_2 reduction.

Light intensity dependence of the reaction quantum yield

In Fig. 10, the initial reaction rates are not proportional to the input power. We reported earlier that the reaction quantum yield (Φ_{CO}) of ZnP-phen=Re significantly decreases as the light intensity increases, due to an inner-filter effect by the OERS of the porphyrin unit which has strong absorption at the excitation wavelength.²³ Fig. 12 shows the plots of Φ_{CO} with respect to the light intensity. The ZnP-*n*Bpy=Re^{Br} series showed similar light intensity dependences. The Φ_{CO} converges under very low light intensity ($<3 \times 10^{-10}$ einstein per s) and finally reaches 24% in ZnP-5Bpy=Re^{Br}. Upon comparing the Φ_{CO} at a low light intensity ($3.2\text{--}3.5 \times 10^{-9}$ einstein per s), we note that the value

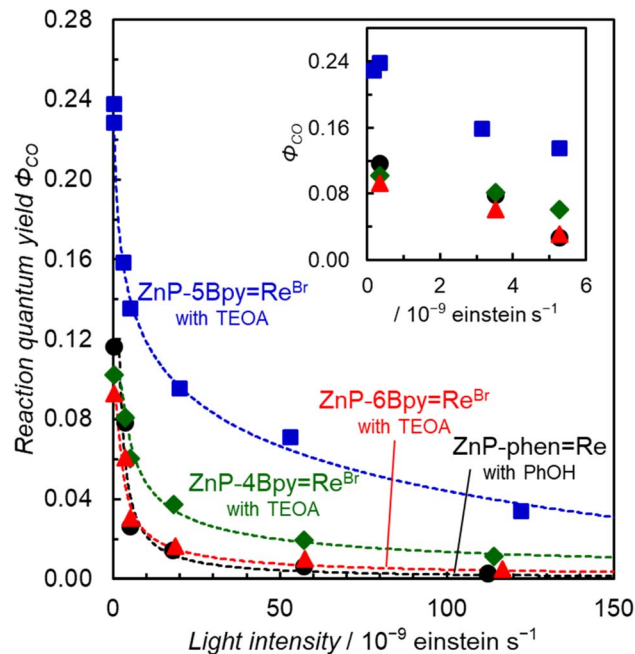


Fig. 12 Dependence of Φ_{CO} on the irradiation light intensity (Xe lamp, $\lambda_{\text{ex}} = 420 \text{ nm}$) for the dyads ZnP-4Bpy=Re^{Br} (green diamond), ZnP-5Bpy=Re^{Br} (blue square), and ZnP-6Bpy=Re^{Br} (red triangle) (0.05 mM) in CO_2 -saturated DMA-TEOA (4 : 1 v/v) solutions containing BIH (0.05 M), and for ZnP-phen=Re (black circle) (0.05 mM) in CO_2 -saturated DMA containing BIH (0.05 M) and phenol (0.1 M). Inset shows magnification of CO formation dependence at low light intensity.

of ZnP-5Bpy=Re^{Br} is twice higher than the values of the other dyads (Table 3).

Fig. 13 shows the UV-vis absorption spectral changes during weak light irradiation at 420 nm. The formations of the OERS of the porphyrin units at *ca.* 450 and 700 nm⁴¹ and chlorins at *ca.* 610 nm⁴² were observed in ZnP-4Bpy=Re^{Br}, ZnP-6Bpy=Re^{Br}, and ZnP-phen=Re (Fig. 13a, c and d). Chlorins are formed by two-electron reduction of porphyrin.^{19c} Indications are therefore that the accumulation of two or more electrons occurs in ZnP-4Bpy=Re^{Br}, ZnP-6Bpy=Re^{Br}, and ZnP-phen=Re. On the other hand, ZnP-5Bpy=Re^{Br} undergoes hardly any change during the irradiation (Fig. 13b). Thus, the reduced species tend to accumulate on the porphyrin in ZnP-4Bpy=Re^{Br}, ZnP-6Bpy=Re^{Br}, and ZnP-phen=Re, whereas ZnP-5Bpy=Re^{Br} showed less electron accumulation, resulting in the low light intensity dependence and the high Φ_{CO} .

Mechanistic insight into the photocatalytic reaction in the presence of TEOA

It has also been reported that the catalysis of CO_2 reduction in the *fac*-Re(bpy)(CO)₃X-type complex (X = Cl⁻, Br⁻, I⁻) commences with the dissociation of a halogen anion upon reduction (Fig. S90[†]).^{2b,43} No induction periods in the time-course profiles of CO formation were observed in the ZnP-*n*Bpy=Re^{Br} series (Fig. 7 and 9), which suggests that efficient reduction of the Re(bpy)(CO)₃Br unit occurs to dissociate the halogen ligand in the three dyads. Considering that the cationic



Table 3 Reaction quantum yields (Φ_{CO}) for the ZnP-*n*Bpy=Re^{Br} series with TEOA and ZnP-phen=Re with phenol^a

Light intensity/einstein per s	ZnP-4Bpy=Re ^{Br}	ZnP-5Bpy=Re ^{Br}	ZnP-6Bpy=Re ^{Br}	ZnP-phen=Re
$3.2\text{--}3.5 \times 10^{-9}$	8%	16%	6%	8%
3×10^{-10}	10%	24%	9%	11%

^a Irradiated at 420 nm (Xe lamp, $\lambda_{\text{ex}} = 420$ nm) to CO₂-saturated DMA-TEOA (4 : 1 v/v) containing the ZnP-*n*Bpy=Re^{Br} series (0.05 mM) and BIH (0.05 M). In ZnP-phen=Re, phenol (0.1 M) was present, not TEOA.

solvent-substituted Re complexes, [Re(bpy)(CO)₃(solvent)]⁺, exhibit more positive reduction potentials than the neutral *fac*-Re(bpy)(CO)₃X-type complexes,⁴³ the reduction of the Re

complex unit is also expected to occur efficiently under the catalytic cycle (Fig. S90[†]). Increasing the concentration of Br ions decreased the catalytic activity (Fig. S91[†]), supporting the

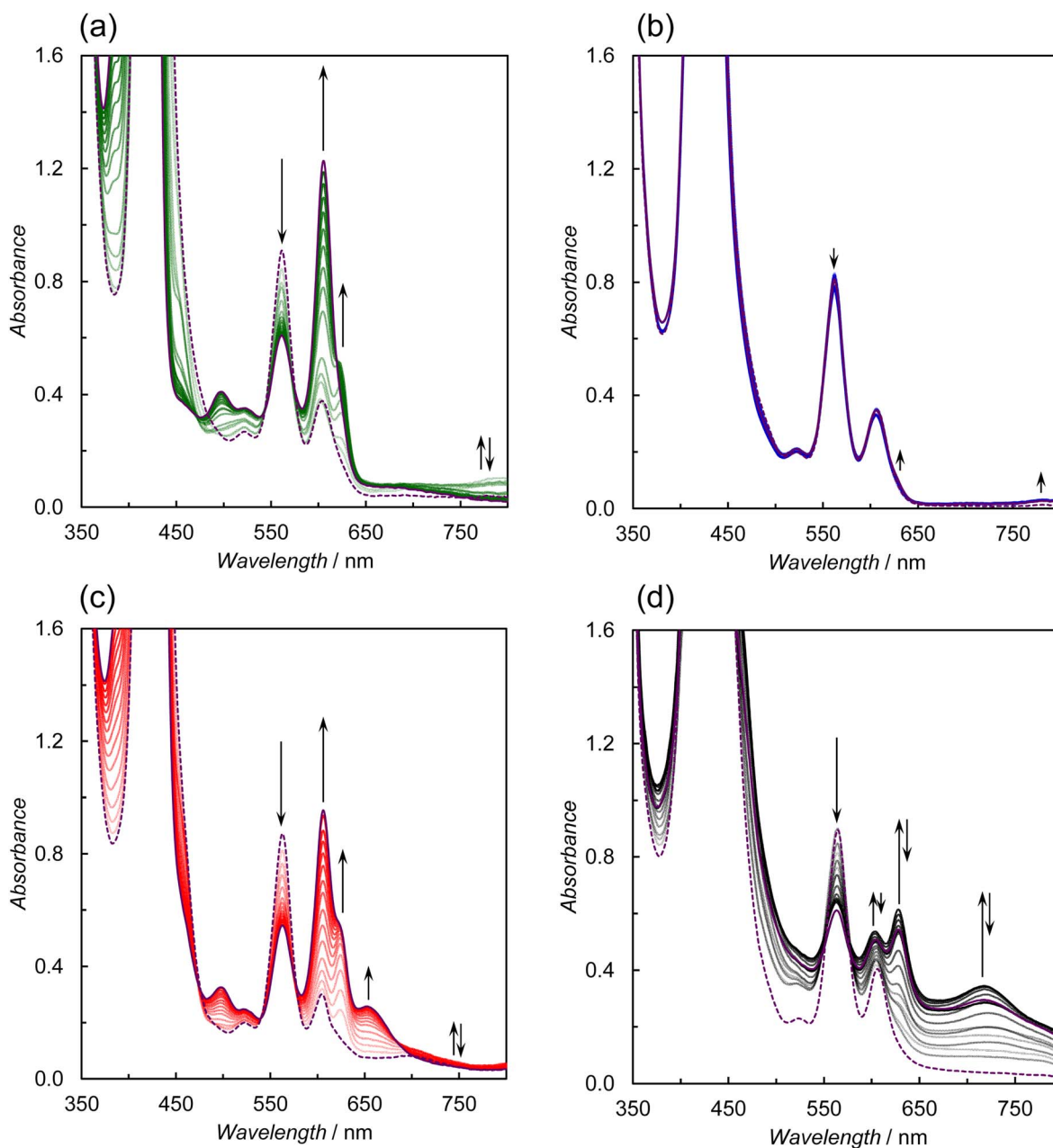


Fig. 13 UV-vis absorption spectral changes during irradiation (0–60 min) at 420 nm (Xe lamp, 5.0×10^{-9} einstein per s) of CO₂ saturated DMA-TEOA (4 : 1 v/v) solutions containing BIH (0.05 M) and the dyad (0.05 mM) (a) ZnP-4Bpy=Re^{Br}, (b) ZnP-5Bpy=Re^{Br}, (c) ZnP-6Bpy=Re^{Br}, and of (d) a CO₂-saturated DMA solution containing BIH (0.05 M) and ZnP-phen=Re (0.05 mM) in the presence of phenol (0.1 M). Purple dotted and solid lines show the spectra before and after the irradiation, respectively.



proposed reaction mechanism that the solvent-substituted species after Br dissociation is the active species.

The Φ_{CO} of **ZnP-5Bpy=Re^{Br}** is double that of the other three dyads in DMA-TEOA (Table 3). Furthermore, despite the high catalytic activity, the UV-vis absorption spectra of **ZnP-5Bpy=Re^{Br}** show almost no change during the photocatalytic reaction (Fig. 13b), indicating that the electron consumption by the CO₂ reduction reaction is accelerated, particularly in **ZnP-5Bpy=Re^{Br}**. Considering that the catalytic activities were similar among the three dyads in the absence of TEOA (Fig. 7), the higher Φ_{CO} in **ZnP-5Bpy=Re^{Br}** would be attributed to TEOA. As mentioned in the Introduction, the ground state of [Re(bpy)(CO)₃(solvent)]⁺ can capture CO₂ in the presence of TEOA (Fig. 1). We calculated the spin densities of the OERS of the **ZnP-*n*Bpy=Re^{Br}** series with and without the bromide ion, and their TEOA and CO₂ adducts (Fig. S79[†]). However, there was no significant difference in the spin density among the three dyads. This suggests that there is another activity-enhancing factor besides TEOA.

From the molecular model optimized by the DFT calculation, TEOA would coordinate both the Re complex and the Zn porphyrin only in **ZnP-5Bpy=Re** (Fig. 14). If TEOA is bound *via* the Zn porphyrin, as shown in Fig. 14, it is expected that CO₂ is quickly captured and the reduction reaction proceeds; then, after the CO₂ reduction reaction it is possible that the next CO₂ is captured and so the reaction proceeds. During the catalytic reaction, it is thought that the bromide ion of the Re complex is substituted by the solvent molecule (Fig. S90[†]),^{2b,38} and the coordination of TEOA to the Re complex occurs in the solvent-substituted complex. Thus, the binding ability of TEOA to the DMA-substituted dyads was investigated to verify the hypothesis of the multidentate coordination (Fig. 14).

The formation of the DMA-substituted dyads, **ZnP-5Bpy=Re^{DMA}** and **ZnP-6Bpy=Re^{DMA}**, was monitored with CO stretching bands of IR spectra (Fig. S65[†]). We confirmed that the substitution reaction with DMA was completed before the addition of TEOA, because TEOA irreversibly reacts with the acetonitrile-substituted Re complex to afford the corresponding iminoester complex (Fig. S92[†]).^{5b} TEOA was added to the resulting DMA solutions: DMA/TEOA = 400 : 1 (v/v). The formation of the **Re-TEOA** structure in Fig. 1 was monitored

with the CO stretching bands' shift to lower wavenumbers in IR spectra (Fig. S93[†]). The CO stretching band at 2000–2050 cm⁻¹ of the thermodynamically equilibrated mixture was separated by curve fitting (Fig. 15). The absorption band profiles were well fitted with two Gaussian-shape functions, corresponding to the DMA-substituted dyad and the TEOA-coordinated dyad (**ZnP-5Bpy=Re^{TEOA}** or **ZnP-6Bpy=Re^{TEOA}**). The absorption intensities as the band areas in **ZnP-5Bpy=Re** and **ZnP-6Bpy=Re** were analyzed as 4 : 6 and 8 : 2 mixtures of the DMA- and TEOA-coordinated dyads, respectively. The equilibrium constants assuming $K = [\text{ZnP-}n\text{Bpy=Re}^{\text{TEOA}}][\text{DMA}]/[\text{ZnP-}n\text{Bpy=Re}^{\text{DMA}}][\text{TEOA}]^2$ were estimated to be 1000 and 110 for **ZnP-5Bpy=Re^{DMA}** and **ZnP-6Bpy=Re^{DMA}**, respectively, indicating the stronger TEOA coordination to the Re complex unit in **ZnP-5Bpy=Re^{DMA}** than in **ZnP-6Bpy=Re^{DMA}**.

We then attempted to verify that the Zn porphyrin unit contributes to the TEOA coordination in **ZnP-5Bpy=Re^{DMA}** through the ¹H NMR spectra in DMF-*d*₇. However, the addition of one equivalent of TEOA hardly showed signal shifts because the binding ability of TEOA to the dyad was too weak in the presence of the competing coordinating DMF. The contribution of the Zn porphyrin unit was verified through UV-vis absorption titration experiments. In general, the Soret and Q-bands of Zn porphyrin are significantly red-shifted by axial coordination of the Zn porphyrin. Upon the addition of TEOA, no spectral change was observed in DMA because DMA coordinates to the Zn porphyrin unit. Thus, TEOA was added to **ZnP-5Bpy=Re^{DMA}** and **ZnP-6Bpy=Re^{DMA}** dissolved in dichloromethane, which is a noncoordinating solvent against Zn porphyrin (Fig. S94[†]).

As control experiments, we also carried out titrations using **ZnP-5Bpy=Re^{Br}** and **ZnP-5Bpy** (Scheme 1). During the titration of TEOA, all the dyads showed red-shifted Soret and Q-bands with isosbestic points (Fig. S95[†]). Spectral change converged with the addition of *ca.* 6 mM TEOA for **ZnP-5Bpy=Re^{DMA}**, but did not converge without the addition of >150 mM TEOA for the other dyads (Fig. S96[†]). The apparent binding constants assuming $K_{\text{app}} = [\text{ZnP-TEOA}]/[\text{ZnP}][\text{TEOA}]$ (ZnP = Zn porphyrin unit) were estimated using Benesi-Hildebrand plots (Fig. S97[†]).⁴⁴ Results showed approximately one order of magnitude larger value for **ZnP-5Bpy=Re^{DMA}** ($K_{\text{app}} = 330 \text{ M}^{-1}$) than for the other three dyads under Ar atmosphere (Table 4).

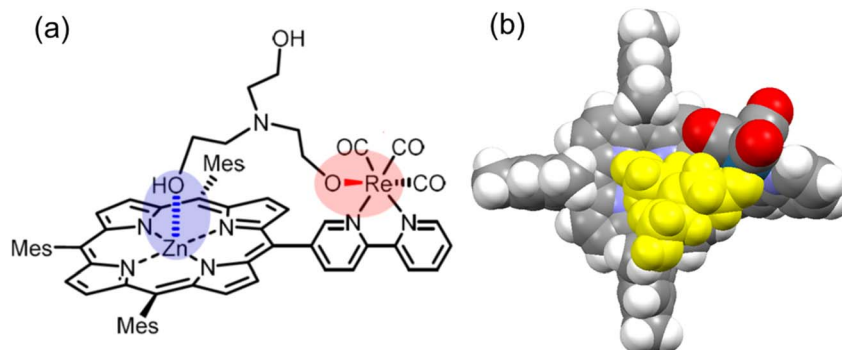


Fig. 14 (a) Plausible structure and (b) molecular model (optimized by DFT calculation) of a TEOA adduct of **ZnP-5Bpy=Re** (without the bromide ion). TEOA is colored in yellow.



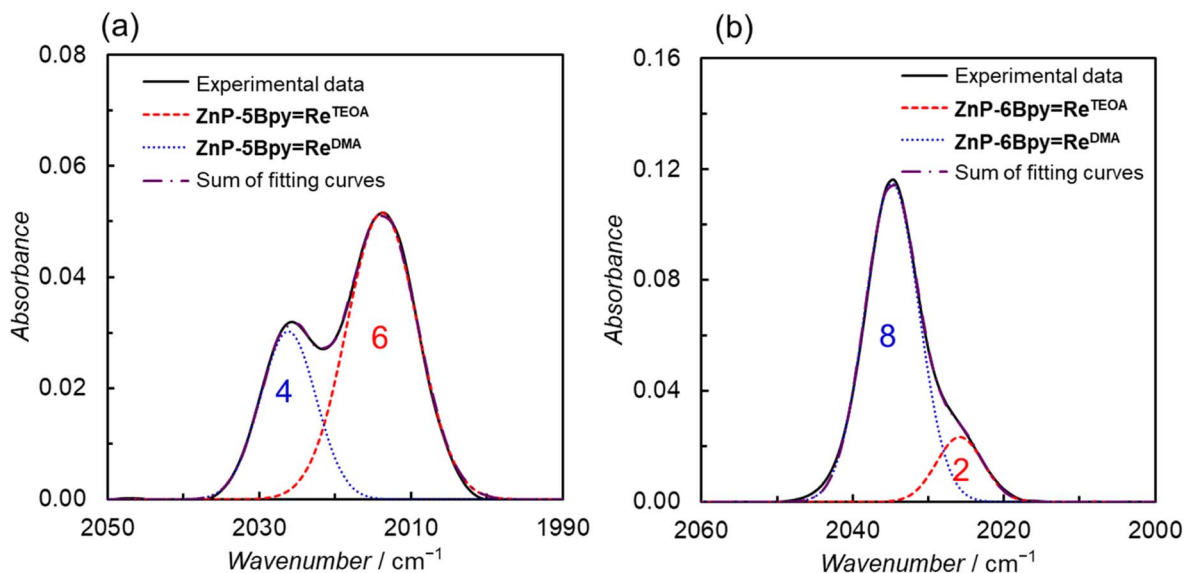


Fig. 15 IR spectra (solid line) of the equilibrated mixture of (a) ZnP-5Bpy=Re^{DMA} and ZnP-5Bpy=Re^{TEOA}, and (b) ZnP-6Bpy=Re^{DMA} and ZnP-6Bpy=Re^{TEOA} in DMA-TEOA (400 : 1 v/v) with the fitting curves (dotted lines). Concentrations of the dyads: 9 mM.

This indicates that after dissociation of the bromide ion a TEOA molecule interacts on both the Re complex and Zn porphyrin units in ZnP-5Bpy=Re^{DMA}, as shown in Fig. 14.

In order to confirm whether TEOA can coordinate properly to the Zn ion even in DMA-TEOA (4 : 1 v/v), the titration experiments with DMA were carried out using ZnP-5Bpy=Re^{Br} and ZnP-5Bpy=Re^{DMA} in dichloromethane (Fig. S98†). Based on the apparent binding constants of DMA coordination, we estimated that 67% of the Zn porphyrin was coordinated with TEOA in Ar-saturated DMA-TEOA (4 : 1 v/v). This suggests that TEOA can sufficiently coordinate with the Zn ion under catalytic reaction conditions. It should be noted that since deprotonation of the OH groups of TEOA is unlikely to occur in dichloromethane, the actual proportion of the TEOA-coordinated Zn porphyrin would be greater than 67% in DMA-TEOA (4 : 1 v/v), where the OH groups of TEOA are more readily deprotonated.

The titration experiments under CO₂ atmosphere for ZnP-5Bpy=Re^{DMA} and ZnP-5Bpy=Re^{Br} showed that the binding constant for ZnP-5Bpy=Re^{DMA} increased significantly to $K_{app} = 760 \text{ M}^{-1}$, compares to that under Ar atmosphere (Table 4 and Fig. S99†). This suggests that the CO₂ insertion into the Re-OR bond occurs, enhancing the binding of TEOA. The binding constant of ZnP-5Bpy=Re^{Br} was also found to be larger under

CO₂ atmosphere than under Ar atmosphere. While it has been reported that CO₂ is captured by insertion into the Zn-methanol bonds,¹⁰ the titration experiments using ZnTMP with TEOA showed similar binding constants ($K_{app} \sim 20 \text{ M}^{-1}$) under both Ar and CO₂ atmospheres. Therefore, it is speculated that the enhanced interaction under CO₂ is mainly due to the interaction around the Re complex.

The IR spectra and results of UV-vis experiments suggest that TEOA strongly binds to the Re complex unit in ZnP-5Bpy=Re^{DMA} by the dual interaction of both the Re complex and the Zn porphyrin units. The tightly bound TEOA would facilitate the CO₂ capture in the ground state of the dyad, resulting in the reaction with CO₂ immediately after the reduction of the Re complex unit to prevent the accumulation of the OERS of the porphyrin (Fig. S100†).⁴⁵ A detailed reaction mechanism involving TEOA and ZnP-5Bpy=Re has not been established yet—further time-resolved spectroscopic measurements are still required.

Conclusions

We systematically synthesized three Zn porphyrin = Re complex dyads, ZnP-4Bpy=Re^{Br}, ZnP-5Bpy=Re^{Br}, and ZnP-6Bpy=Re^{Br}, to investigate the effect of the connecting positions between the Re complex and the Zn porphyrin units on photophysical properties and the catalytic activity of CO₂ reduction. Although RTP was observed only in ZnP-6Bpy=Re^{Br}, there was no significant difference in the catalytic activity and durability of the three dyads when only BIH was used (as electron donor) without additives. The durability of the present three dyads was inferior to that of the previously reported dyad ZnP-phen=Re as a result of the dissociation of the Re complex moiety during the reduction reaction. The addition of TEOA suppressed the Re dissociation and remarkably improved the activity and

Table 4 Apparent binding constants^a in dichloromethane from UV-vis absorption spectra

	K_{app}/M^{-1} , under Ar	K_{app}/M^{-1} , under CO ₂
ZnP-5Bpy	31	—
ZnP-5Bpy=Re ^{Br}	30	48
ZnP-5Bpy=Re ^{DMA}	330	760
ZnP-6Bpy=Re ^{DMA}	48	—

^a $K_{app} = [\text{ZnP-TEOA}]/[\text{ZnP}][\text{TEOA}]$.



durability in the CO₂ reduction reaction. The initial reaction rate of **ZnP-5Bpy=Re^{Br}** was unexpectedly more than twice that of the other dyads. The reaction quantum yield of **ZnP-5Bpy=Re^{Br}** reached 24%—the highest yet recorded for the photocatalytic CO₂ reduction using porphyrins as photosensitizers. The higher Φ_{CO} value originated from no electron accumulation at the porphyrin unit during the photocatalytic reaction as well as acceleration of the CO₂ reduction. The IR and UV-vis titration experiments suggested a dual interaction of TEOA *via* the Zn porphyrin and the Re complex units in **ZnP-5Bpy=Re^{DMA}**, allowing a stronger interaction of TEOA to the Re complex. Although **ZnP-5Bpy=Re^{Br}** exhibited no RTP, the dependence of Φ_{CO} on the BIH concentration suggests that the catalytic reaction is mediated *via* the T₁ of the Zn porphyrin unit. This therefore indicates that the RTP is not an essential property for porphyrin to act as a good triplet photosensitizer for the photocatalytic CO₂ reduction.

We reconfirmed that, when using porphyrin as photosensitizer, the balance of the electron relay cycle on the photosensitizer and the catalytic cycle on the catalyst^{2b} is crucial to achieving high durability for CO₂ reduction. The prompt catalytic reaction, with the assistance of the coordination of TEOA, plays an important role in the high durability of the photosensitizer. In particular, a dual interaction *via* the Zn porphyrin further promotes the catalytic reaction. The enhancement of the coordination of TEOA has been also observed in other metal complex catalysts.^{6–11} The present dual interaction of TEOA *via* a coordination scaffold suggests new opportunities to achieve enhanced catalytic performance by capturing CO₂ with high efficiency, and subsequently the further development of practical photocatalytic systems using a wide range of organic dyes that include porphyrin-based photosensitizers.

Experimental section

General procedure

All chemicals and solvents were of commercial reagent quality and were used without further purification unless otherwise stated. Syntheses of *fac*-Re(bpy)(CO)₃Br and BIH were carried out according to the literature.^{3a,12} DMA and TEOA were dried over molecular sieves of size 4A and distilled under reduced pressure. Dry toluene was prepared by distillation over benzophenone/Na. Toluene used for fluorescence quantum yields and lifetimes was dealt with conc. H₂SO₄ after simple distillation, and washed with H₂O and Na₂CO₃ aq. After dried over CaCl₂, the toluene was distilled on CaH₂. Dry dichloromethane was dried over molecular sieves of size 4A. Pyrrole was prepared by distillation over CaH₂. The reactions were monitored on silica-gel 60 F254 TLC plates (Merck). Silica-gels utilized for column chromatography were purchased from Kanto Chemical Co. Inc.: Silica-Gel 60N (Spherical, Neutral) 63–210 μm and (Flash) 40–50 μm . Alumina used for column chromatography were purchased from Merck (aluminum oxide 90 standardized). ¹H and ¹³C NMR, distortionless enhancement by polarization transfer 135 (DEPT 135), ¹H–¹H COSY spectra were recorded by using JEOL JNM-ECS300 (300 MHz), JNM-ECZ400S (400 MHz), JNM-ECA500II (500 MHz), and Bruker AVANCE NEO

400 (400 MHz). Chemical shifts were recorded in parts per million (ppm) relative to tetramethylsilane. UV-vis absorption spectra were collected using square cells (optical path length = 10 mm) on JASCO V-650 and V-660 spectrometers. Steady-state emission spectra were collected on Hitachi F-4500 spectrofluorometer or Horiba FluoroMAX 4 and corrected for the response of the detector system. The fluorescence intensities were normalized at the absorbance of the excitation wavelength. Fluorescence quantum yields were determined by integrated ratios of the fluorescence spectra relative to that of ZnTPP (Φ_{f} = 3.3% in toluene).⁴⁶ Their values were corrected by refractive indexes of the used solvents. Time-resolved emission signals for fluorescence lifetimes were obtained by a pico-second fluorescence measurement system consisting of Chromex 250IS imaging spectrograph and a charge-coupled device (CCD) streak scope Hamamatsu model C4334, with excitation by a light pulser Hamamatsu model PLP-1-040 (406 nm, full width at half-maximum (fwhm) = 70 ps).⁴⁷ Hi-resolution MALDI-TOF mass spectra were collected on JEOL JMS S-3000 with dithranol or *trans*-2-[3-(4-*tert*-butylphenyl)-2-methyl-2-propenylidene]-malononitrile (DCTB) as a matrix. FTIR spectra were recorded in KBr or acetonitrile using a JASCO FT/IR-4600. CV and DPV were measured using an ALS-H/CHI Model 612E electrochemical analyzer (BAS) in a micro-cell equipped with a glassy carbon working electrode (ϕ 1.6 mm) and a Pt counter-electrode. The micro-cell was connected *via* a Luggin capillary with a reference electrode of Ag/AgNO₃ (10 mM in DMA). Tetrabutylammonium hexafluorophosphate (ⁿBu₄NPF₆) recrystallized from ethyl acetate was used as a supporting electrolyte. Ferrocene was used as an external standard, and all potentials were referenced to the ferrocene/ferrocenium couple ($E_{1/2}$ = +0.099 V vs. Ag/AgNO₃). UV-vis spectral changes during irradiation were obtained using an Asahi Spectra PRA-201 apparatus equipped with a Xe lamp (Asahi Spectra, MAX-350).

5,10,15-Trimesityl-20-(2-bromopyridin-6-yl)-21H,23H-porphyrin (H₂P-6py)

In a 1 L flask were placed 2,4,6-trimethylbenzaldehyde (542 mg, 3.7×10^{-3} mol), 6-bromo-2-pyridinecarboxaldehyde (226 mg, 1.2×10^{-3} mol), pyrrole (327 mg, 4.9×10^{-3} mol), and CHCl₃ (500 mL). The solution was degassed by bubbling with Ar gas, and BF₃OEt₂ (270 mg, 1.9×10^{-3} mol) was added. After the reaction mixture was stirred for 1 h at room temperature, triethylamine (370 mg, 3.7×10^{-3} mol) and *p*-chloranil (900 mg, 3.7×10^{-3} mol) were added and stirring was continued for 19 h. The crude product obtained by evaporation of the solvent was purified with flash silica gel columns (CHCl₃:hexane = 1:1). The second band was collected and the solvent was evaporated to dryness, giving the title compound as a purple solid (171 mg, 17% based on 6-bromo-2-pyridinecarboxaldehyde). TLC (silica gel, CHCl₃) R_{f} = 0.6, (alumina, CHCl₃) R_{f} = 0.8; MALDI-TOF-mass (dithranol) m/z [M + H]⁺ 820.3019, calcd for [C₅₂H₄₇BrN₅]⁺ 820.3009; ¹H NMR (400 MHz, CDCl₃) δ /ppm = 8.78 (d, J = 4.7 Hz, 2H, β -pyrrole), 8.71 (d, J = 4.7 Hz, 2H, β -pyrrole), 8.64 (d, J = 4.7 Hz, 2H, β -pyrrole), 8.62 (d, J = 4.7 Hz, 2H, β -pyrrole), 8.08 (dd, J = 4.7 and 3.6 Hz, 1H, py), 7.90 (d, J =



4.7 Hz, 1H, py), 7.89 (d, $J = 3.6$ Hz, 1H, py), 7.25 (s, 6H, mesityl), 2.62 (s, 9H, CH₃), 1.84 (s, 18H, CH₃), -2.58 (s, 2H, NH); ¹³C NMR (100 MHz, CDCl₃) δ /ppm = 161.94 (C), 141.02 (C), 139.54 (C), 138.36 (C), 138.17 (C), 137.88 (C), 136.95 (CH), 130.65 (CH), 129.55 (CH), 127.89 (CH), 126.96 (CH), 118.95 (C), 118.47 (C), 115.19 (C), 21.80 (CH), 21.59 (CH).

5,10,15-Trimesityl-20-(2-bromopyridin-5-yl)-21H,23H-porphyrin (H₂P-5py)

In a 2 L flask were placed 2,4,6-trimethylbenzaldehyde (1.9 g, 13×10^{-3} mol), 6-bromo-3-pyridinecarboxaldehyde (800 mg, 4.3×10^{-3} mol), pyrrole (1.2 g, 17×10^{-3} mol), and CHCl₃ (1.7 L). The solution was degassed by bubbling with N₂ gas, and BF₃OEt₂ (1.3 mL, 11×10^{-3} mol) diluted with CHCl₃ (50 mL) was added through a dropping funnel over 4 h. After the reaction mixture was further stirred for 5 h at room temperature, triethylamine (2.1 mL, 15×10^{-3} mol) and *p*-chloranil (3.2 g, 13×10^{-3} mol) were added and stirring was continued for 20 h. The resulting solution was filtered through a Celite pad and the filtrate was evaporated to dryness *in vacuo*. The crude product was purified with an alumina column (CHCl₃), and silica gel columns twice (CHCl₃:hexane = 2:1) and flash silica gel columns twice (CHCl₃:hexane = 2:1 → CHCl₃), giving the title compound as a purple solid (380 mg, 14% based on 6-bromo-3-pyridinecarboxaldehyde). TLC (silica gel, CHCl₃) $R_f = 0.6$, (alumina, CHCl₃) $R_f = 0.8$; MALDI-TOF-mass (dithranol) m/z [M + H]⁺ 820.2969, calcd for [C₅₂H₄₇BrN₅]⁺ 820.3009; ¹H NMR (400 MHz, CDCl₃) δ /ppm = 9.20 (d, $J = 2.5$ Hz, 1H, py), 8.76 (d, $J = 4.7$ Hz, 2H, β -pyrrole), 8.71 (d, $J = 4.7$ Hz, 2H, β -pyrrole), 8.67 (s, 4H, β -pyrrole), 8.31 (dd, $J = 2.5$ and 8.0 Hz, 1H, py), 7.85 (d, $J = 8.0$ Hz, 1H, py), 7.28 (s, 4H, mesityl), 7.27 (s, 2H, mesityl), 2.61 (s, 6H, CH₃), 2.60 (s, 3H, CH₃), 1.86 (s, 18H, CH₃), -2.55 (s, 2H, NH); ¹³C NMR (100 MHz, CDCl₃) δ /ppm = 153.74 (CH), 143.30 (CH), 142.08 (C), 139.63 (C), 139.59 (C), 139.55 (C), 138.30 (C), 138.15 (C), 138.09 (C), 138.06 (C), 137.65 (C), 130.71 (CH), 128.09 (CH), 128.03 (CH), 126.46 (CH), 118.91 (C), 118.77 (C), 112.87 (C), 21.98 (C), 21.92 (C), 21.68 (C).

5,10,15-Trimesityl-20-(2-bromopyridin-4-yl)-21H,23H-porphyrin (H₂P-4py)

In a 1 L flask were placed 2,4,6-trimethylbenzaldehyde (880 mg, 5.9×10^{-3} mol), 2-bromo-4-pyridinecarboxaldehyde (370 mg, 2.0×10^{-3} mol), pyrrole (530 mg, 7.9×10^{-3} mol), and CHCl₃ (790 mL). The solution was degassed by bubbling with Ar gas, and BF₃OEt₂ (550 mg, 3.9×10^{-3} mol) was added. After the reaction mixture was stirred for 3 h at room temperature, triethylamine (600 mg, 5.9×10^{-3} mol) and *p*-chloranil (1.5 g, 5.9×10^{-3} mol) were added and stirring was continued for 3 h. The crude product obtained by evaporation of the solvent was purified with alumina columns twice (CHCl₃) and a flash silica gel column (CHCl₃:hexane = 1:1 → CHCl₃), giving the title compound as a purple solid (156 mg, 10% based on 2-bromo-4-pyridinecarboxaldehyde). TLC (silica gel, CHCl₃) $R_f = 0.5$, (alumina, CHCl₃) $R_f = 0.8$; MALDI-TOF-mass (dithranol) m/z [M + H]⁺ 820.2981, calcd for [C₅₂H₄₇BrN₅]⁺ 820.3009; ¹H NMR (400 MHz, CDCl₃) δ /ppm = 8.75 (d, $J = 4.8$ Hz, 3H, β -pyrrole and py),

8.72 (d, $J = 4.8$ Hz, 2H, β -pyrrole), 8.66 (s, 4H, β -pyrrole), 8.35 (d, $J = 1.3$ Hz, 1H, py), 8.11 (dd, $J = 1.3$ and 4.8 Hz, 1H, py), 7.28 (s, 4H, mesityl), 7.27 (s, 2H, mesityl), 2.62 (s, 6H, CH₃), 2.61 (s, 3H, CH₃), 1.86 (s, 18H, CH₃), -2.60 (s, 2H, NH); ¹³C NMR (100 MHz, CDCl₃) δ /ppm = 153.55 (C), 148.28 (CH), 141.40 (C), 139.49 (C), 138.13 (C), 138.06 (C), 138.02 (C), 133.32 (CH), 130.66 (CH), 128.63 (CH), 127.98 (CH), 127.94 (CH), 119.13 (C), 118.82 (C), 113.55 (C), 21.89 (CH), 21.81 (CH), 21.60 (CH).

Zinc(II) 5,10,15-trimesityl-20-(2-bromopyridin-6-yl)-21H,23H-porphyrin (ZnP-6py)

A saturated methanolic solution of zinc(II) acetate (270 mg, 1.5×10^{-3} mol) was added to a solution of H₂P-6py (120 mg, 1.5×10^{-4} mol) in CHCl₃ (30 mL). The mixture was stirred overnight at room temperature. The resulting solution was diluted with CHCl₃ and washed with water. The organic layer was dried over anhydrous Na₂SO₄ and evaporated to dryness. The further purification was carried out with a flash silica gel column (CHCl₃:hexane = 1:1), giving the titled compound as a purple solid (130 mg, 95%). TLC (silica gel, CHCl₃) $R_f = 0.5$; MALDI-TOF-mass (dithranol) m/z [M]⁺ 881.1969, calcd for [C₅₂H₄₄BrN₅Zn]⁺ 881.2072; ¹H NMR (400 MHz, CDCl₃) δ /ppm = 8.83 (d, $J = 4.7$ Hz, 2H, β -pyrrole), 8.79 (d, $J = 4.7$ Hz, 2H, β -pyrrole), 8.72 (d, $J = 4.6$ Hz, 2H, β -pyrrole), 8.70 (d, $J = 4.6$ Hz, 2H, β -pyrrole), 8.20 (dd, $J = 1.5$ and 6.8 Hz, 1H, py), 7.93 (dd, $J = 6.8$ and 8.0 Hz, 1H, py), 7.89 (dd, $J = 1.5$ and 8.0 Hz, 1H, py), 7.27 (s, 6H, mesityl), 2.63 (s, 9H, CH₃), 1.96 (s, 3H, CH₃), 1.83 (s, 15H, CH₃); ¹³C NMR (100 MHz, CDCl₃) δ /ppm = 162.81 (C), 150.28 (C), 149.86 (C), 149.83 (C), 149.55 (C), 140.76 (C), 139.40 (C), 139.03 (C), 138.98 (C), 137.56 (C), 136.80 (CH), 131.51 (CH), 131.46 (CH), 131.34 (CH), 131.18 (CH), 128.97 (CH), 127.76 (CH), 126.77 (CH), 119.75 (C), 119.30 (C), 116.08 (C), 21.77 (CH), 21.57 (CH).

Zinc(II) 5,10,15-trimesityl-20-(2-bromopyridin-5-yl)-21H,23H-porphyrin (ZnP-5py)

In a 200 mL flask, a methanolic solution saturated with zinc(II) acetate (2.2 g, 12×10^{-3} mol) was added to a solution of H₂P-5py (490 mg, 6.0×10^{-4} mol) in CHCl₃ (120 mL). The mixture was stirred overnight at room temperature. The resulting solution was diluted with CHCl₃ and washed with water. The organic layer was dried over anhydrous Na₂SO₄ and evaporated to dryness. The further purification was carried out with a flash silica gel column (CHCl₃:hexane = 1:2). The collected fraction was evaporated, giving the titled compound as a purple solid (430 mg, 81%). TLC (silica gel, CHCl₃) $R_f = 0.5$; MALDI-TOF-mass (dithranol) m/z [M]⁺ 881.2034, calcd for [C₅₂H₄₄BrN₅Zn]⁺ 881.2072; ¹H NMR (400 MHz, CDCl₃) δ /ppm = 9.01 (s, 1H, py), 8.80 (d, $J = 4.4$ Hz, 2H, β -pyrrole), 8.72 (m, 6H, β -pyrrole), 8.33 (d, $J = 7.9$ Hz, 1H, py), 7.85 (d, $J = 7.9$ Hz, 1H, py), 7.27 (s, 6H, mesityl), 2.63 (s, 9H, CH₃), 1.87 (s, 3H, CH₃), 1.83 (s, 15H, CH₃); ¹³C NMR (100 MHz, CDCl₃ + 1 drop of CD₃OD) δ /ppm = 153.31 (CH), 150.15 (C), 149.96 (C), 149.70 (C), 149.14 (C), 143.30 (CH), 139.39 (C), 139.32 (C), 139.28 (C), 138.98 (C), 137.35 (C), 131.20 (CH), 131.07 (CH), 131.03 (CH), 130.69 (CH), 127.60 (CH), 126.04 (CH), 119.03 (C), 118.85 (C), 21.71 (CH), 21.43 (CH).



Zinc(II) 5,10,15-trimesityl-20-(2-bromopyridin-4-yl)-21H,23H-porphyrin (ZnP-4py)

A saturated methanolic solution of zinc(II) acetate (350 mg, 1.9×10^{-3} mol) was added to a solution of **H₂P-4py** (160 mg, 1.9×10^{-4} mol) in CHCl₃ (38 mL). The mixture was stirred overnight at room temperature. The resulting solution was diluted with CHCl₃ and washed with water. The organic layer was dried over anhydrous Na₂SO₄ and evaporated to dryness. The further purification was carried out with a flash silica gel column (CHCl₃ : hexane = 1 : 1), giving the titled compound as a purple solid (170 mg, 99%). TLC (silica gel, CHCl₃) *R_f* = 0.4; MALDI-TOF-mass (dithranol) *m/z* [M]⁺ 881.2040, calcd for [C₅₂H₄₄BrN₅Zn]⁺ 881.2072; ¹H NMR (400 MHz, CDCl₃) δ/ppm = 8.81 (d, *J* = 4.7 Hz, 2H, β-pyrrole), 8.75 (d, *J* = 4.7 Hz, 2H, β-pyrrole), 8.72 (d, *J* = 4.7 Hz, 2H, β-pyrrole), 8.71 (d, *J* = 4.7 Hz, 2H, β-pyrrole), 8.55 (dd, *J* = 5.0, 0.7 Hz, 1H, py), 8.32 (dd, *J* = 1.5, 0.7 Hz, 1H, py), 8.09 (dd, *J* = 5.0, 1.5 Hz, 1H, py), 7.28 (s, 4H, mesityl), 7.27 (s, 2H, mesityl), 2.63 (s, 6H, CH₃), 2.62 (s, 3H, CH₃), 1.85 (s, 3H, CH₃), 1.84 (s, 3H, CH₃), 1.83 (s, 6H, CH₃), 1.82 (s, 6H, CH₃); ¹³C NMR (100 MHz, CDCl₃) δ/ppm = 154.38 (C), 150.40 (C), 150.17 (C), 149.89 (C), 148.42 (C), 147.69 (CH), 140.77 (C), 139.31 (C), 139.28 (C), 138.88 (C), 137.69 (C), 137.65 (C), 133.32 (C), 131.67 (CH), 131.55 (CH), 131.47 (CH), 130.96 (CH), 128.46 (CH), 127.81 (CH), 119.81 (C), 119.56 (C), 114.37 (C), 21.89 (CH), 21.79 (CH), 21.57 (CH).

Zinc(II) 5,10,15-trimesityl-20-(2,2'-bipyridin-6-yl)-21H,23H-porphyrin (ZnP-6Bpy)

In a 200 mL Schlenk tube were placed **ZnP-6py** (150 mg, 1.7×10^{-4} mol), 2-(tributylstannyl)pyridine (120 μL, 3.4×10^{-4} mol), Pd(PPh₃)₄ (42 mg, 3.6×10^{-5} mol), and dry toluene (57 mL). The reaction mixture was degassed by freeze–thaw cycles and stirred at 105 °C overnight. The solvent of the resulting mixture was evaporated after addition of water (50 mL). The residue was dissolved in CHCl₃ and washed with water, and the organic layer was dried over anhydrous Na₂SO₄. The crude product obtained by evaporation of the solvent was purified with a flash silica gel column (CHCl₃) and an anhydrous K₂CO₃/flash silica gel (1 : 9 w/w) column (CHCl₃) to remove organotin impurities.³² Further purification was carried out by reprecipitation with CHCl₃/hexane, giving the titled compound as a purple solid (98 mg, 64%). TLC (silica gel, CHCl₃) *R_f* = 0.2, (silica gel, CHCl₃ : acetone = 4 : 1) *R_f* = 0.4; MALDI-TOF-mass (dithranol) *m/z* [M + H]⁺ 881.3202, calcd for [C₅₇H₄₉N₆Zn]⁺ 881.3305; ¹H NMR (500 MHz, CDCl₃) δ/ppm = 8.92 (d, *J* = 4.6 Hz, 2H, β-pyrrole), 8.80 (dd, *J* = 1.1 and 7.7 Hz, 1H, bpy), 8.79 (d, *J* = 4.9 Hz, 1H, bpy), 8.78 (d, *J* = 4.6 Hz, 2H, β-pyrrole), 8.73 (d, *J* = 4.6 Hz, 2H, β-pyrrole), 8.71 (d, *J* = 4.6 Hz, 2H, β-pyrrole), 8.58 (d, *J* = 7.7 Hz, 1H, bpy), 8.25 (dd, *J* = 1.1 and 7.7 Hz, 1H, bpy), 8.20 (t, *J* = 7.7 Hz, 1H, bpy), 7.71 (td, *J* = 7.7 and 1.1 Hz, 1H, bpy), 7.32 (ddd, *J* = 1.1, 4.9 and 7.7 Hz, 1H, bpy), 7.26 (s, 6H, mesityl), 2.63 (s, 3H, CH₃), 2.62 (s, 6H, CH₃), 1.93 (s, 3H, CH₃), 1.88 (s, 6H, CH₃), 1.79 (s, 9H, CH₃); ¹³C NMR (125 MHz, CDCl₃) δ/ppm = 160.99 (C), 156.67 (C), 154.81 (C), 150.14 (C), 149.94 (C), 149.82 (C), 149.31 (CH), 139.42 (C), 139.35 (C), 139.31 (C), 139.12 (C), 139.05 (C), 137.49 (C), 137.09 (CH), 135.52 (CH), 131.85 (CH), 131.32

(CH), 131.20 (CH), 131.10 (CH), 130.17 (CH), 127.72 (CH), 123.91 (CH), 121.99 (CH), 119.45 (CH), 119.31 (C), 119.04 (C), 118.23 (C), 21.78 (CH), 21.55 (CH).

Zinc(II) 5,10,15-trimesityl-20-(2,2'-bipyridin-5-yl)-21H,23H-porphyrin (ZnP-5Bpy)

In a 100 mL Schlenk tube were placed **ZnP-5py** (95 mg, 1.1×10^{-4} mol), 2-(tributylstannyl)pyridine (72 μL, 2.2×10^{-4} mol), Pd(PPh₃)₄ (25 mg, 2.2×10^{-5} mol), and dry toluene (36 mL). The reaction mixture was degassed by freeze–thaw cycles and stirred at 105 °C overnight. The solvent of the resulting mixture was evaporated after addition of water (40 mL). The residue was dissolved in CHCl₃ and washed with water, and the organic layer was dried over anhydrous Na₂SO₄. The crude product obtained by evaporation of the solvent was purified with a flash silica gel column (CHCl₃) and an anhydrous K₂CO₃/flash silica gel (1 : 9 w/w) column (CHCl₃) to remove organotin impurities.³² Further purification was carried out by reprecipitation with CHCl₃/hexane, giving the titled compound as a purple solid (61 mg, 64%). TLC (silica gel, CHCl₃) *R_f* = 0.2, (silica gel, CHCl₃ : acetone = 4 : 1) *R_f* = 0.4; MALDI-TOF-mass (dithranol) *m/z* [M + H]⁺ 881.3335, calcd for [C₅₇H₄₉N₆Zn]⁺ 881.3305; ¹H NMR (400 MHz, CDCl₃) δ/ppm = 9.47 (s, 1H, bpy), 8.87 (d, *J* = 4.6 Hz, 2H, β-pyrrole), 8.82–8.79 (m, 2H, bpy), 8.80 (d, *J* = 4.6 Hz, 2H, β-pyrrole), 8.72 (s, 4H, β-pyrrole), 8.68 (d, *J* = 7.4 Hz, 1H, bpy), 8.66 (d, *J* = 7.4 Hz, 1H, bpy), 7.96 (t, *J* = 7.4 Hz, 1H, bpy), 7.43 (t, *J* = 7.4 Hz, 1H, bpy), 7.27 (s, 6H, mesityl), 2.62 (s, 9H, CH₃), 1.87 (s, 3H, CH₃), 1.85 (s, 3H, CH₃), 1.84 (s, 12H, CH₃); ¹³C NMR (100 MHz, CDCl₃) δ/ppm = 156.38 (C), 154.96 (C), 153.15 (CH), 150.18 (C), 150.07 (C), 149.89 (C), 149.78 (CH), 149.52 (C), 141.97 (CH), 139.38 (C), 139.34 (C), 139.15 (C), 139.09 (C), 137.53 (C), 137.25 (CH), 136.83 (C), 131.58 (CH), 131.41 (CH), 131.31 (CH), 131.20 (CH), 128.83 (CH), 127.75 (CH), 127.01 (CH), 123.98 (CH), 121.56 (CH), 119.18 (CH), 115.41 (C), 21.87 (CH₃), 21.79 (CH₃), 21.55 (CH₃).

Zinc(II) 5,10,15-trimesityl-20-(2,2'-bipyridin-4-yl)-21H,23H-porphyrin (ZnP-4Bpy)

In a 100 mL Schlenk tube were placed **ZnP-4py** (100 mg, 1.2×10^{-4} mol), 2-(tributylstannyl)pyridine (77 μL, 2.3×10^{-4} mol), Pd(PPh₃)₄ (28 mg, 2.4×10^{-5} mol), and dry toluene (39 mL). The reaction mixture was degassed by freeze–thaw cycles and stirred at 105 °C overnight. The solvent of the resulting mixture was evaporated after addition of water (40 mL). The residue was dissolved in CHCl₃ and washed with water, and the organic layer was dried over anhydrous Na₂SO₄. The crude product obtained by evaporation of the solvent was purified with a flash silica gel column (CHCl₃) and an anhydrous K₂CO₃/flash silica gel (1 : 9 w/w) column (CHCl₃) to remove organotin impurities.³² Further purification was carried out by reprecipitation with CHCl₃/hexane, giving the titled compound as a purple solid (44 mg, 43%). TLC (silica gel, CHCl₃) *R_f* = 0.2, (silica gel, CHCl₃ : acetone = 4 : 1) *R_f* = 0.3; MALDI-TOF-mass (dithranol) *m/z* [M + H]⁺ 881.3333, calcd for [C₅₇H₄₉N₆Zn]⁺ 881.3305; ¹H NMR (400 MHz, CDCl₃) δ/ppm = 9.27 (s, 1H, bpy), 9.03 (d, *J* = 4.8 Hz, 1H, bpy), 8.84 (d, *J* = 5.0 Hz, 2H, β-pyrrole), 8.75 (d, *J* = 5.0 Hz, 2H, β-



pyrrole), 8.71 (d, 1H, bpy), 8.70 (s, 4H, β -pyrrole), 8.63 (d, $J = 4.8$ Hz, 1H, bpy), 8.17 (d, $J = 4.8$ Hz, 1H, bpy), 7.92 (t, $J = 7.8$ Hz, 1H, bpy), 7.33 (dd, $J = 7.8, 4.8$ Hz, 1H, bpy), 7.27 (s, 6H, mesityl), 2.62 (s, 9H, CH₃), 1.84 (m, 18H, CH₃); ¹³C NMR (100 MHz, CDCl₃) δ /ppm = 156.56 (C), 154.34 (C), 152.82 (C), 150.11 (C), 149.90 (CH), 149.87 (CH), 149.41 (C), 148.92 (C), 147.36 (CH), 143.24 (CH), 139.39 (C), 139.34 (C), 139.30 (C), 137.42 (C), 137.16 (CH), 131.51 (CH), 131.27 (CH), 131.11 (CH), 131.07 (CH), 129.58 (CH), 127.71 (CH), 126.58 (CH), 123.93 (CH), 121.99 (CH), 121.87 (CH), 119.01 (C), 118.95 (C), 116.54 (C), 21.93 (CH₃), 21.86 (CH₃), 21.82 (CH₃), 21.56 (CH₃).

Dyad connected *via* the 6-position of the Bpy ligand (ZnP-6Bpy=Re^{Br})

In a 200 mL Schlenk tube were placed ZnP-6Bpy (49 mg, 5.5×10^{-5} mol) and bromopentacarbonylrhenium(i) (23 mg, 5.5×10^{-5} mol), and toluene (37 mL). The reaction mixture was stirred at 90–95 °C overnight. The solvent of the resulting mixture was evaporated and the residue was purified with a flash silica gel column (CHCl₃ → CHCl₃: acetone = 4:1). Further purification was carried out by recrystallization with CHCl₃/hexane followed by CHCl₃/ethanol, giving the titled compound as needle purple crystals (32 mg, 47%). TLC (silica gel, CHCl₃) $R_f = 0.1$, (silica gel, CHCl₃: acetone = 4:1) $R_f = 0.6$; MALDI-TOF-mass (DCTB) m/z [M]⁺ 1228.1670, calcd for [C₆₀H₄₈BrN₆O₃-ReZn]⁺ 1228.1792; ¹H NMR (400 MHz, CDCl₃) δ /ppm = 9.01 (d, $J = 5.5$ Hz, 1H, bpy), 8.99 (d, $J = 4.6$ Hz, 1H, β -pyrrole), 8.81 (d, $J = 4.6$ Hz, 1H, β -pyrrole), 8.72 (d, $J = 4.6$ Hz, 1H, β -pyrrole), 8.71 (s, 2H, β -pyrrole), 8.70 (d, $J = 4.6$ Hz, 1H, β -pyrrole), 8.68 (d, $J = 4.6$ Hz, 1H, β -pyrrole), 8.66 (d, $J = 7.8$ Hz, 1H, bpy), 8.53 (d, $J = 8.0$ Hz, 1H, bpy), 8.48 (d, $J = 7.8$ Hz, 1H, bpy), 8.32 (t, $J = 7.8$ Hz, 1H, bpy), 8.29 (d, $J = 4.6$ Hz, 1H, β -pyrrole), 8.16 (td, $J = 8.0, 1.8$ Hz, 1H, bpy), 7.51 (dd, $J = 8.0, 5.5$ Hz, bpy), 7.29 (s, 1H, mesityl), 7.26 (s, 3H, mesityl), 7.24 (s, 1H, mesityl), 7.22 (s, 1H, mesityl), 2.63 (s, 3H, CH₃), 2.61 (s, 3H, CH₃), 2.60 (s, 3H, CH₃), 1.97 (s, 3H, CH₃), 1.94 (s, 3H, CH₃), 1.89 (s, 3H, CH₃), 1.79 (s, 3H, CH₃), 1.76 (s, 3H, CH₃), 1.74 (s, 3H, CH₃); ¹H NMR (500 MHz, DMF-*d*₇) δ /ppm = 9.16 (m, 1H, bpy), 9.04 (d, $J = 8.0$ Hz, 1H, bpy), 8.93 (dd, $J = 5.4, 1.8$ Hz, 1H, bpy), 8.84 (d, $J = 4.6$ Hz, 1H, β -pyrrole), 8.55–8.44 (m, 7H, β -pyrrole and bpy $\times 2$), 8.43 (d, $J = 4.6$ Hz, 1H, β -pyrrole), 8.35 (td, $J = 8.0, 1.8$ Hz, 1H, bpy), 8.32 (d, $J = 4.6$ Hz, 1H, β -pyrrole), 7.68 (ddd, $J = 8.0, 5.4, 1.6$ Hz, 1H, bpy), 7.24 (s, 1H, mesityl), 7.21 (s, 1H, mesityl), 7.20 (s, 1H, mesityl), 7.19 (s, 1H, mesityl), 7.17 (s, 2H, mesityl), 2.47 (s, 3H, CH₃), 2.46 (s, 3H, CH₃), 2.43 (s, 3H, CH₃), 1.88 (s, 3H, CH₃), 1.71 (s, 3H, CH₃), 1.70 (s, 3H, CH₃), 1.68 (s, 3H, CH₃), 1.66 (s, 3H), 1.56 (s, 3H, CH₃); ¹³C NMR (125 MHz, DMF-*d*₇) δ /ppm = 199.03 (CO), 191.93 (CO), 189.13 (CO), 164.82 (C), 158.29 (C), 155.73 (C), 153.09 (CH), 150.30 (C), 150.20 (C), 149.92 (C), 149.83 (C), 149.73 (C), 149.38 (C), 149.29 (C), 149.27 (C), 148.09 (CH), 140.04 (CH), 139.67 (C), 139.61 (C), 139.43 (C), 139.25 (C), 138.98 (C), 138.76 (C), 138.68 (C), 138.59 (C), 137.44 (C), 137.38 (C), 137.34 (CH), 136.44 (CH), 132.54 (CH), 132.09 (CH), 131.22 (CH), 130.95 (CH), 130.83 (CH), 130.62 (CH), 130.55 (CH), 130.44 (CH), 127.84 (CH), 127.79 (CH), 127.73 (CH), 127.66 (CH), 127.30 (CH), 125.66 (CH), 123.71 (CH), 123.64 (CH), 119.36 (C), 118.81

(C), 118.76 (C), 117.54 (C), 21.45 (CH₃), 21.26 (CH₃), 21.16 (CH₃), 21.06 (CH₃), 20.85 (CH₃), 20.81 (CH₃); FT-IR (KBr) $\nu_{CO} = 1895, 1923, 2021$ cm⁻¹, (acetonitrile) $\nu_{CO} = 1894, 1919, 2021$ cm⁻¹.

Dyad connected *via* the 5-position of the Bpy ligand (ZnP-5Bpy=Re^{Br})

In a 200 mL Schlenk tube were placed ZnP-5Bpy (57 mg, 6.4×10^{-5} mol) and bromopentacarbonylrhenium(i) (26 mg, 6.5×10^{-5} mol), and toluene (43 mL). The reaction mixture was stirred at 90–95 °C overnight. The solvent of the resulting mixture was evaporated and the residue was recrystallized with CHCl₃/hexane followed by CHCl₃/ethanol, giving the titled compound as needle brown crystals (44 mg, 55%). TLC (silica gel, CHCl₃) $R_f = 0.2$, (silica gel, CHCl₃: acetone = 4:1) $R_f = 0.7$; MALDI-TOF-mass (DCTB) m/z [M]⁺ 1228.1768, calcd for [C₆₀H₄₈BrN₆O₃-ReZn]⁺ 1228.1792; ¹H NMR (400 MHz, CDCl₃) δ /ppm = 9.89 (s, 1H, bpy), 9.20 (d, $J = 5.7$ Hz, 1H, bpy), 8.93 (d, $J = 4.9$ Hz, 1H, β -pyrrole), 8.89 (d, $J = 4.9$ Hz, 2H, β -pyrrole), 8.81 (m, 2H, β -pyrrole, bpy), 8.75 (s, 4H, β -pyrrole), 8.54 (d, $J = 8.3$ Hz, 1H, bpy), 8.45 (d, $J = 8.1$ Hz, 1H, bpy), 8.15 (dd, $J = 8.1, 7.8$ Hz, 1H, bpy), 7.60 (dd, $J = 7.8, 5.7$ Hz, 1H, bpy), 7.27 (m, 6H, mesityl), 2.62 (s, 9H, CH₃), 1.88 (s, 3H, CH₃), 1.87 (s, 3H, CH₃), 1.85 (s, 9H, CH₃), 1.80 (s, 3H, CH₃); ¹³C NMR (100 MHz, CDCl₃) δ /ppm = 197.06 (CO), 196.55 (CO), 189.21 (CO), 158.60 (C), 156.57 (CH), 156.15 (C), 154.51 (CH), 154.06 (C), 153.60 (CH), 150.77 (C), 150.52 (C), 150.32 (C), 149.75 (C), 149.29 (C), 148.79 (C), 143.22 (CH), 139.49 (C), 139.44 (C), 139.20 (C), 139.07 (C), 138.86 (C), 138.82 (C), 138.75 (C), 138.65 (C), 138.65 (C), 137.75 (C), 132.66 (CH), 132.08 (CH), 131.92 (CH), 131.86 (CH), 131.70 (CH), 131.58 (CH), 131.38 (CH), 130.32 (CH), 128.02 (CH), 127.93 (CH), 127.87 (CH), 127.77 (CH), 127.15 (CH), 125.98 (CH), 125.82 (CH), 123.64 (CH), 121.18 (CH), 120.33 (C), 120.09 (C), 119.92 (C), 111.25 (C), 21.92 (CH₃), 21.83 (CH₃), 21.75 (CH₃), 21.57 (CH₃); FT-IR (KBr) $\nu_{CO} = 1900, 1921, 2022$ cm⁻¹, (acetonitrile) $\nu_{CO} = 1902, 1918, 2023$ cm⁻¹.

Dyad connected *via* the 4-position of the Bpy ligand (ZnP-4Bpy=Re^{Br})

In a 200 mL Schlenk tube were placed ZnP-4Bpy (44 mg, 5.0×10^{-5} mol) and bromopentacarbonylrhenium(i) (20 mg, 5.0×10^{-5} mol), and toluene (33 mL). The reaction mixture was stirred at 90–95 °C overnight. The solvent of the resulting mixture was evaporated and the residue was purified with flash silica gel columns (CHCl₃ → CHCl₃: acetone = 4:1). Further purification was carried out by recrystallization with CHCl₃/hexane followed by CHCl₃/ethanol, giving the titled compound as needle dark-green crystals (25 mg, 41%). TLC (silica gel, CHCl₃) $R_f = 0.3$, (silica gel, CHCl₃: acetone = 4:1) $R_f = 0.8$; MALDI-TOF-mass (DCTB) m/z [M]⁺ 1228.1757, calcd for [C₆₀H₄₈BrN₆O₃-ReZn]⁺ 1228.1792; ¹H NMR (500 MHz, CDCl₃) δ /ppm = 9.43 (d, $J = 5.6$ Hz, 1H, bpy), 9.24 (d, $J = 5.6$ Hz, 1H, bpy), 9.03 (d, $J = 1.7$ Hz, 1H, bpy), 8.87 (d, $J = 4.6$ Hz, 1H, β -pyrrole), 8.83 (d, $J = 4.6$ Hz, 2H, β -pyrrole), 8.80 (d, $J = 4.6$ Hz, 1H, β -pyrrole), 8.75–8.72 (m, 4H, β -pyrrole), 8.38 (dd, $J = 5.6, 1.7$ Hz, 1H, bpy), 8.29 (d, $J = 8.3$ Hz, 1H, bpy), 8.02 (t, $J = 8.3$ Hz, 1H, bpy), 7.59 (dd, $J = 8.3, 5.6$ Hz, 1H, bpy), 7.29 (s, 2H, mesityl), 7.28 (s, 2H, mesityl),



7.26 (s, 2H, mesityl), 2.63 (s, 9H, CH₃), 1.87 (s, 3H, CH₃), 1.85 (s, 3H, CH₃), 1.84 (s, 9H, CH₃), 1.82 (s, 3H, CH₃); ¹³C NMR (125 MHz, CDCl₃) δ/ppm = 196.98 (CO), 196.94 (CO), 189.36 (CO), 156.33 (C), 155.24 (C), 153.75 (C), 153.68 (CH), 151.27 (CH), 150.77 (C), 150.60 (C), 150.33 (C), 150.13 (C), 149.79 (C), 148.20 (C), 147.96 (C), 139.33 (C), 139.24 (C), 139.16 (C), 139.08 (C), 138.93 (CH), 137.87 (C), 138.65 (C), 138.57 (C), 137.98 (C), 137.87 (C), 137.83 (C), 137.78 (C), 132.29 (CH), 132.21 (CH), 132.03 (CH), 131.95 (CH), 131.73 (CH), 131.63 (CH), 130.95 (CH), 130.29 (CH), 129.13 (CH), 128.47 (CH), 128.33 (CH), 127.98 (CH), 127.92 (CH), 127.88 (CH), 127.83 (CH), 127.20 (CH), 125.39 (CH), 123.51 (CH), 120.48 (C), 120.16 (C), 119.97 (C), 113.20 (C), 21.91 (CH₃), 21.86 (CH₃), 21.80 (CH₃), 21.57 (CH₃); FT-IR (KBr) ν_{CO} = 1902, 1920, 2021 cm⁻¹.

Acetonitrile-substituted ZnP-6Bpy=Re (ZnP-6Bpy=Re^{MeCN})

In a 50 mL flask were placed ZnP-6Bpy=Re^{Br} (10 mg, 8.1 × 10⁻⁶ mol), AgBF₄ (1.9 mg, 9.7 × 10⁻⁶ mol), and acetonitrile (4 mL). The reaction mixture was stirred at reflux for 4 h, and then the solvent was evaporated to dryness. The crude product was dissolved in CHCl₃ (50 mL) and washed with water (50 mL × 3). The organic layer was passed through Phase Separator paper (Whatman). After evaporating the solvent, the residue was purified with a flash silica gel column (CHCl₃ → CHCl₃: acetonitrile = 10 : 1), giving a purple-green solid (18 mg, 90%). TLC (silica gel, CHCl₃: acetonitrile = 10 : 1) R_f = 0.3; ¹H NMR (400 MHz, CDCl₃) δ/ppm = 9.02 (dd, J = 8.2, 1.4 Hz, 1H, bpy), 8.89 (d, J = 8.2, 1.1 Hz, 1H, bpy), 8.86–8.84 (m, 1H, bpy), 8.83 (d, J = 4.6 Hz, 1H, β-pyrrole), 8.79 (d, J = 4.6 Hz, 1H, β-pyrrole), 8.77–8.71 (m, 5H, β-pyrrole), 8.62 (dd, J = 8.2, 1.4 Hz, 1H, bpy), 8.54 (t, J = 8.2 Hz, 1H, bpy), 8.50 (d, J = 4.6 Hz, 1H, β-pyrrole), 8.36 (ddd, J = 8.2, 7.6, 1.4 Hz, 1H, bpy), 7.62 (ddd, J = 7.6, 5.4, 1.4 Hz, 1H, bpy), 7.32 (s, 1H, mesityl), 7.30 (s, 2H, mesityl), 7.28 (m, 3H, mesityl), 2.65 (s, 3H, CH₃), 2.64 (s, 3H, CH₃), 2.64 (s, 3H, CH₃), 2.34 (s, 3H, coordinated acetonitrile), 2.01 (s, 3H, CH₃), 1.88 (s, 3H, CH₃), 1.86 (s, 6H, CH₃), 1.84 (s, 3H, CH₃), 1.80 (s, 3H, CH₃); ¹⁹F NMR (375 MHz, CDCl₃) δ/ppm = -152 (BF₄⁻); FT-IR (acetonitrile) ν_{CO} = 1921, 1941, 2034 cm⁻¹.

Acetonitrile-substituted ZnP-5Bpy=Re (ZnP-5Bpy=Re^{MeCN})

In a 50 mL flask were placed ZnP-5Bpy=Re^{Br} (7.0 mg, 5.7 × 10⁻⁶ mol), AgBF₄ (1.3 mg, 6.8 × 10⁻⁶ mol), and acetonitrile (4 mL). The reaction mixture was stirred at reflux for 4 h, and then the solvent was evaporated to dryness. The crude product was dissolved in CHCl₃ (50 mL) and washed with water (50 mL × 3). The organic layer was passed through Phase Separator paper (Whatman). After evaporating the solvent, the residue was purified with a flash silica gel column (CHCl₃ → CHCl₃: acetonitrile = 10 : 1), giving a purple-green solid (12 mg, 86%). TLC (silica gel, CHCl₃: acetonitrile = 10 : 1) R_f = 0.3; ¹H NMR (400 MHz, CDCl₃) δ/ppm = 9.70 (s, 1H, bpy), 9.07 (d, J = 5.2 Hz, 1H, bpy), 9.04–8.98 (m, 2H, bpy and β-pyrrole), 8.97–8.87 (m, 3H, bpy and β-pyrrole), 8.81 (d, J = 8.8 Hz, 1H, bpy), 8.79–8.70 (m, 5H, β-pyrrole), 8.38 (t, J = 8.0 Hz, 1H, bpy), 7.73 (t, J = 6.5 Hz, 1H, bpy), 7.30 (s, 1H, mesityl), 7.28 (s, 5H, mesityl), 2.63 (s, 9H, CH₃), 2.42 (s, 3H, coordinated acetonitrile), 1.90 (s, 3H, CH₃),

1.87 (s, 3H, CH₃), 1.85 (s, 6H, CH₃), 1.83 (s, 3H, CH₃), 1.81 (s, 3H, CH₃); ¹⁹F NMR (375 MHz, CDCl₃) δ/ppm = -152 (BF₄⁻); FT-IR (acetonitrile) ν_{CO} = 1926, 1940, 2037 cm⁻¹.

Quenching experiments

The Stern–Volmer relationship (eqn (1)) was obtained by the plots of the relative emission band intensity (*I*₀/*I*) versus the concentration of BIH:

$$I_0/I = 1 + K_{SV}[\text{BIH}] = 1 + k_q\tau[\text{BIH}] \quad (1)$$

where *I*₀ and *I* represent the emission intensities in the absence and the presence of an electron donor (quencher), BIH, respectively. *K*_{SV}, *k*_q, τ are the Stern–Volmer constant, the quenching rate constant, and an emission lifetime, respectively. The Ar-saturated DMA solutions containing the porphyrins in square quartz cells (optical path length = 10 mm) were bubbled with Ar gas and the emission spectra of each samples were measured.

Photocatalytic CO₂ reduction

In glass tubes (8.0 mL, i.d. = 10 mm), 1.4 mL of CO₂-saturated solutions containing BIH was added by 0.6 mL of Ar-saturated solutions containing the photocatalysts, and the reaction solutions were bubbled through septum caps with CO₂ gas for 20 min. Photo-irradiations to determine TON were carried out using a merry-go-round irradiation apparatus (Iris-MG, Cell Systems) equipped with LED lamps at 298 K. Reaction quantum yields (Φ_{CO}) and UV-vis absorption spectral changes during photoirradiation were determined with an Asahi Spectra PRA-201 apparatus equipped with a Xe lamp (Asahi Spectra, MAX-350). The irradiation of very weak light (<5 × 10⁻¹⁰ einstein per s) was carried out with a Hitachi F-4500 spectrophotometer and the number of the incident photons was determined by a standard chemical actinometer using K₃[Fe(C₂O₄)₃].⁴⁸ Reaction quantum yields were calculated according to Φ_{CO} = (amounts of CO)/(incident photons). In 7 mL quartz cubic cells (7.0 mL, optical path length = 10 mm), 2.4 mL of CO₂-saturated solutions containing BIH was added by 0.6 mL of Ar-saturated solutions containing the photocatalysts, and the reaction solutions were bubbled through septum caps with CO₂ gas for 20 min. The gaseous reaction products (CO and H₂) were quantified with a gas chromatography system (GC-2014, Shimadzu Science) equipped with a Shincarbon column (i.d. 3.0 mm × 3.0 m, Shinwa Chemical Industries) and a thermal conductivity detector (TCD). The product (HCOOH) in the solutions was analyzed with a capillary electrophoresis system (Otsuka Electronics Co. CAPI-3300I).

Computational methods

DFT calculations were carried out using the Gaussian 09 package of programs.⁴⁹ Each structure was fully optimized using the B3LYP functional using the 6-31G(d) basis set for all atoms except Re and the standard double-ζ type LANL2DZ basis set with the effective core potential of Hay–Wadt for Re. The stationary points were verified using the vibrational analysis.



The calculations were carried out by using the polarizable continuum model (PCM) with default parameter for DMA.

Equilibrium constants of TEOA in DMA

$\text{ZnP-}n\text{Bpy}=\text{Re}^{\text{MeCN}}$ ($n = 5, 6$) was added to Ar-saturated DMA (dyad concentration 11 mM) and the ligand substitutions from acetonitrile to DMA were monitored by IR spectroscopy. An Ar-saturated DMA-TEOA (79 : 1 v/v) solution (60 μL) was added to equilibrated DMA solutions of the dyads (240 μL) and the resulting mixtures were bubbled with Ar gas for 10 min. The IR spectra were monitored. The concentrations of $\text{ZnP-}n\text{Bpy}=\text{Re}^{\text{DMA}}$ and $\text{ZnP-}n\text{Bpy}=\text{Re}^{\text{TEOA}}$ were estimated based on the deconvolution of the absorption bands in Fig. 15, which were obtained by curve fittings as a linear combination of the Gaussian functions with OriginPro software. The equilibrium constants in DMA were defined as $K = [\text{ZnP-}n\text{Bpy}=\text{Re}^{\text{TEOA}}]/[\text{DMA}][\text{ZnP-}n\text{Bpy}=\text{Re}^{\text{DMA}}][\text{TEOA}]$.⁵

Apparent binding constants of TEOA in dichloromethane

Ar-saturated DMA solutions containing $\text{ZnP-}n\text{Bpy}=\text{Re}^{\text{MeCN}}$ ($n = 5, 6$) were allowed to stand for 2 h prior to collecting samples of solutions (20 μL) and drying *in vacuo*. Solid samples were dissolved in Ar- or CO_2 -saturated dichloromethane to prepare dichloromethane solutions containing the dyads (3 μM). Solutions of $\text{ZnP-}5\text{Bpy}=\text{Re}^{\text{Br}}$ and $\text{ZnP-}5\text{Bpy}$ were prepared by dissolving in Ar- or CO_2 -saturated dichloromethane to be 3 μM . Titrations were carried out in square cells (optical path length 10 mm) by adding three dichloromethane solutions containing TEOA (0.1, 1.0, and 10 M) *via* septum caps. The apparent binding constants of TEOA to the Zn porphyrins in dichloromethane were defined as $K_{\text{app}} = [\text{TEOA-coordinated Zn porphyrin}]/[\text{Zn porphyrin}][\text{TEOA}]$. The values of K_{app} were estimated from spectral data using Benesi-Hildebrand plot assuming the formation of a 1 : 1 complex of TEOA and Zn porphyrins (Fig. S96[†]).

Time-resolved near-infrared emission measurements

Near-infrared time-resolved emission spectra and lifetimes were measured by a lab-made system controlled by a Labview-based program. Samples were excited at 406 nm pulses with an appropriate duration time (100 ns to 100 μs) which were prepared by combination of an ORBIS 405LX laser and a Stanford Research Systems (SRS) digital pulse generator Model DG535. After dispersed by a Horiba spectrometer MicroHR, emission signals were detected on a Hamamatsu R5509-43 photomultiplier by the single-photon-counting method. SRS Multichannel scaler SRS430 was used for signal counting and accumulation. Two dimensional time-wavelength data sets were acquired to a PC and emission spectra for an appropriate window were reconstructed from the 2D dataset.

Data availability

All the data supporting this article have been included in the ESI.[†]

Author contributions

Y. K. designed the dyads, conceived the idea of the dual interaction of TEOA, performed a part of the titration experiments, and wrote the manuscript. Y. S. synthesized and characterized the dyads, and performed most of photocatalytic experiments. S. A. and T. S. performed the time-resolved emission measurements. H. I. contributed to the synthesis of a portion of the dyads. M. S. A. supervised the time-resolved emission measurements and analyzed the data, and provided suggestions on writing the manuscript. A. S. supervised the project, and provided suggestions on the experiments and writing the manuscript.

Conflicts of interest

There are no conflicts to declare.

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